

TISSUE FACTOR ANTAGONIST AND PROTEIN C POLYPEPTIDE COMPOSITIONS

## CROSS-REFERENCE TO RELATED APPLICATIONS

This application claims priority under 35 U.S.C. 119 of Danish application no. PA 2002 5 01709 filed November 6, 2002 and U.S. application no. 60/434,911 filed December 20, 2002, the contents of which are fully incorporated herein by reference.

## FIELD OF THIS INVENTION

The invention relates to pharmaceutical compositions comprising a Tissue Factor (TF) 10 antagonist and protein C, protein C-related polypeptide, or combination thereof, methods of using such compositions or combinations of TF antagonist and protein C and/or protein C-related peptide(s) to induce, promote, and/or enhance a desired physiological effect in a subject, and other compositions and methods related to such compositions, combinations, and methods.

## 15 BACKGROUND OF THE INVENTION

Tissue Factor (TF) is a cellular transmembrane receptor for plasma coagulation factor VIIa (FVIIa). The formation of TF/VIIa complexes on the cell surface is believed to trigger the coagulation cascade *in vivo*. The TF/VIIa complex efficiently activates coagulation factor IX and factor X. The resultant protease factor Xa (Xa), converts prothrombin to thrombin, which in turn 20 converts fibrinogen into a fibrin matrix. Normally, TF is constitutively expressed on the surface of extravascular cells that are not in contact with the blood, but is not expressed on the surface of cells that come in contact with blood. TF is also expressed in various pathophysiological conditions where it is believed to be involved in progression of disease states within cancer, inflammation, atherosclerosis, and ischemia/reperfusion.

25 Recently, TF has been shown to also act as a mediator of intracellular activities either by interactions of the cytoplasmic domain of TF with the cytoskeleton or by supporting the VIIa-protease dependent signaling. Such activities may be responsible, at least partly, for the implicated role of TF in tumor development, metastasis, and angiogenesis. Cellular exposure of TF activity is advantageous in a crisis of vascular damage but may be fatal when exposure is 30 sustained (as it is in these and other disease states).

Factor VIIa (FVIIa) is a two-chain, 50 kilodalton (kDa) vitamin-K dependent, plasma serine protease which participates in the complex regulation of *in vivo* hemostasis. FVIIa is generated from proteolysis of a single peptide bond from its single chain zymogen, Factor VII (FVII). The conversion of FVII into FVIIa occurs by cleavage of an internal peptide bond. In the 35 presence of calcium ions, FVIIa binds with high affinity to exposed TF, which acts as a cofactor for FVIIa, enhancing the proteolytic activation of its substrates FVII, Factor IX, and FX. Inactivated FVII (FVIIai) is FVIIa modified in such a way that the protein has been rendered catalytically

inactive. FVIIai is thus not able to catalyze the conversion of FX to FXa, but still able to bind to TF in competition with active endogenous FVIIa and thereby inhibit TF activity.

Activated Protein C (APC) is a serine protease and naturally occurring anticoagulant that plays a role in the regulation of vascular homeostasis by inactivating Factors Va and VIIIa in 5 the coagulation cascade. Human Protein C (PC) is made in vivo primarily in the liver as a single polypeptide of 461 amino acids. In concert with other proteins, Protein C functions as an important down-regulator of blood coagulation factors that promote thrombosis.

It is often desirable to selectively block the coagulation cascade in a patient. Anticoagulants such as heparin, coumarin, derivatives of coumarin, indandione derivatives, or 10 other agents may be used, for example, during kidney dialysis, or to treat deep vein thrombosis, disseminated intravascular coagulation (DIC), and in the treatment of similar disorders. Treatment with heparin and other known anticoagulants may, however, have undesirable side 15 effects. For example, available anticoagulants generally act throughout the body, rather than acting specifically at a clot site. Heparin, which currently is widely used, may cause heavy bleeding. Moreover, because heparin acts as a cofactor for antithrombin III (AT III), and AT III is rapidly depleted in DIC treatment, it is often difficult to maintain the proper heparin dosage, necessitating continuous monitoring of AT III and heparin levels. Heparin is also ineffective if AT 20 III depletion is extreme. Further, prolonged use of heparin may also increase platelet aggregation and reduce platelet count, and has been implicated in the development of osteoporosis. Heparin is not the only currently used anticoagulant with serious potential side effects. Indandione derivatives, for example, may have toxic side effects.

Inhibitors of tissue factor may act as antagonists for tissue factor-mediated induction of coagulation and, thus, block the production of thrombin and the subsequent deposition of fibrin. The use of FVIIai compositions has been proposed as an inhibitor of TF. For example, 25 International Patent Applications WO 92/15686, WO 94/27631, WO 96/12800, WO 97/47651 disclose FVIIai compositions and the uses thereof. International Patent Applications WO 90/03390, WO 95/00541, WO 96/18653, and European Patent 500 800 describe peptides derived from FVIIa that have TF/FVIIa antagonist activity. International patent application WO 01/21661 discloses a bivalent inhibitor of FVII and FXa. Other proposed strategies include using 30 monoclonal antibodies (Mabs), catalytically impaired FVIIa mutants and chemically inactivated FVIIa, as TF antagonists. For example, mouse Mabs against TF are described in US Patents 6,001,978 and 5,223,427, International Patent Application WO 99/51743 discloses human/mouse chimeric monoclonal antibodies directed against human TF, European Patent Application 833 911 describes CDR-grafted antibodies against human TF, and Presta et al., *Thrombosis and 35 Haemostasis*, 85(3):379-389 (2001) describes a humanized antibody against TF. Despite the description of these compositions, no TF antagonists have been developed and marketed for

therapeutic use in humans. Moreover, the applicability of such compositions and strategies may not be suited to all uses and/or all patients.

Accordingly, there remains a need for alternative and improved compositions having anticoagulant and anti-inflammatory activity. There is especially a need for compositions that 5 can induce, promote, and/or enhance such activities when administered at relatively low doses, and that desirably not produce the undesirable side effects associated with traditional anticoagulant compositions. The invention described herein fulfils these needs by providing anticoagulants that act specifically at sites of injury or TF exposure, and further provides other related advantages. Furthermore the invention described herein provides compounds, which 10 acts to inhibit the cellular functions of TF, which can be useful in the treatment and/or prevention of conditions like sepsis, inflammation, atherosclerosis, restenosis, and/or cancer. These and other advantages, as well as additional inventive features, will be apparent from the description of the invention provided herein.

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#### SUMMARY OF THE INVENTION

The invention described herein provides novel and useful compositions comprising a Tissue Factor (TF) antagonist and protein C, protein C-related polypeptide, or combination thereof. The invention also provides novel and useful methods of using such compositions or combinations of TF antagonist and protein C and/or protein C-related peptide(s) to induce, 20 promote, and/or enhance a desired physiological effect in a subject. Other features of the invention arise from compositions and methods related to such compositions, combinations, and methods, described herein.

In one exemplary aspect, the invention provides a composition comprising (a) a pharmaceutically acceptable carrier, diluent, and/or excipient, (b) a tissue factor antagonist, and 25 (c) protein C or a protein C-related polypeptide.

In another exemplary aspect, the invention provides a method of inducing, promoting, and/or enhancing at least one physiological response associated with the prevention or treatment of a thrombotic disease, coagulopathic disease, respiratory disease, or inflammatory disease associated with TF in a subject suffering from or at risk of acquiring such a disease 30 comprising administering a TF antagonist and a protein C or a protein C-related polypeptide to the subject in an amount sufficient to detectably induce, promote, and/or enhance the physiological response.

These and other aspects and features of the invention are more fully described in the following detailed description of the invention.

## DETAILED DESCRIPTION OF THIS INVENTION

As described above, the invention described herein provides a composition comprising a TF antagonist and protein C or a protein C-related polypeptide. The composition can be a pharmaceutically acceptable composition and/or can comprise one or more pharmaceutically acceptable carriers, diluents, and/or excipients, and the like. The composition can include or be free from any additional number of active and/or inactive agents and, thus, can consist or consist essentially of a preparation of a TF antagonist and a preparation of protein C or a protein C-related polypeptide, and, optionally, one or more of the components selected from pharmaceutical acceptable excipients, carriers, stabilizers, detergents, neutral salts, antioxidants, preservatives, and protease inhibitors.

In one aspect, the composition is in single-dosage form. In another aspect, the composition is in the form of a first-unit dosage form and a second-unit dosage form, where the first-unit dosage form comprises, consists, or consists essentially of a preparation of a TF antagonist and one or more of the components selected from the list of pharmaceutical acceptable excipients or carriers, stabilizers, detergents, neutral salts, antioxidants, preservatives, and protease inhibitors; and the second-unit dosage form comprises, consists, or consists essentially of a preparation of protein C or a protein C-related polypeptide and one or more of the components selected from the list of pharmaceutical acceptable excipients or carriers, stabilizers, detergents, neutral salts, antioxidants, preservatives, and protease inhibitors.

Unless otherwise stated, the TF antagonist can be any suitable antagonist. In a particular aspect, the TF antagonist is or comprises a factor VII polypeptide that has been chemically inactivated in the active site. Such a TF antagonist can be modified by any suitable method so as to inactivate the active site. For example, the TF antagonist can be or comprise a factor VII polypeptide that is inactivated in the active site by reaction with a reagent selected from peptide chloromethylketones or peptidyl chloromethanes; azapeptides; acylating agents such as various guanidinobenzoate derivatives and 3-alkoxy-4-chloroisocoumarins; sulphonyl fluorides such as phenylmethylsulphonylfluoride (PMSF); diisopropylfluorophosphate (DFP); tosylpropylchloromethyl ketone (TPCK); tosylsylchloromethyl ketone (TLCK); nitrophenylsulphonates; heterocyclic protease inhibitors such as isocoumarines and coumarins; and any suitable combinations thereof. The TF antagonist also can be inactivated in the active site by reaction with a reagent selected from the list of: Phe-Phe-Arg chloromethyl ketone, Phe-Phe-Arg chloromethylketone, D-Phe-Phe-Arg chloromethyl ketone, D-Phe-Phe-Arg chloromethylketone Phe-Pro-Arg chloromethylketone, D-Phe-Pro-Arg chloromethylketone, Phe-Pro-Arg chloromethylketone, D-Phe-Pro-Arg chloromethylketone, L-Glu-Gly-Arg chloromethylketone and D-Glu-Gly-Arg chloromethylketone, Dansyl-Phe-Phe-Arg chloromethyl ketone, Dansyl-Phe-Phe-Arg chloromethylketone, Dansyl-D-Phe-Phe-Arg chloromethyl ketone, Dansyl-D-Phe-Phe-Arg chloromethylketone, Dansyl-Phe-Pro-Arg chloromethylketone, Dansyl-D-

Phe-Pro-Arg chloromethylketone, Dansyl-Phe-Pro-Arg chloromethylketone, Dansyl-D-Phe-Pro-Arg chloromethylketone, Dansyl-L-Glu-Gly-Arg chloromethylketone, Dansyl-D-Glu-Gly-Arg chloromethylketone, and any suitable combination thereof.

The TF antagonist also can be or can comprise an antibody against TF. The TF antibody 5 can be any suitable antibody, such as a monoclonal antibody, for example, a fully human monoclonal antibody. In other particular aspects, the TF antagonist is a humanized monoclonal antibody, such as a mouse/human chimeric antibody. In one aspect, the antibody is an antibody against human TF. In other particular aspects, the TF antagonist is or comprises an antibody selected from a Fab fragment; a monovalent fragment consisting of the VL, VH, CL and CH I 10 domains; a F(ab)2 fragment; a F(ab')2 fragment; a bivalent fragment comprising two Fab fragments linked by a disulfide bridge at the hinge region; a Fd fragment consisting of the VH and CH1 domains; a Fv fragment consisting of the VL and VH domains of a single arm of an antibody, a dAb fragment; an isolated complementarity determining region (CDR); or a single chain Fv (scFv). The TF antagonist also can be or comprise a derivative and/or variant of such an 15 antibody or fragment (e.g., the TF antagonist can be an antibody modified by addition of one or more chemical moieties to the amino acid sequence thereof or the addition, substitution, or deletion of one or more amino acid sequences).

In another aspect of the invention, the TF antagonist comprises, consists essentially, or 20 consists of a biologically active Factor VII variant. In other particular aspects of the invention, the TF antagonist is a variant of a naturally occurring Factor VII, e.g., human and bovine Factor VII, wherein the active site residue, (e.g., Ser344), is modified or replaced with Gly, Met, Thr, or more 25 preferably, Ala. Such substitution can be made separately or in combination with substitution(s) at other sites in the catalytic triad of the polypeptide (e.g., His193 and Asp242). The TF antagonist also can be a peptide having an amino acid sequence that comprises or consists essentially of the amino acid sequences of these proteins or biologically active fragments thereof. In other aspects, the TF antagonist is or comprises a variant human factor VII that 30 comprises an amino acid substitution of the lysine corresponding to position 341 of SEQ ID NO: 1. The TF antagonist also or additionally can be or comprise a variant of human Factor VII that comprises an amino acid substitution of the serine corresponding to position 344 of SEQ ID NO: 1. 35 1. The TF antagonist also can be or comprise a human factor VII variant that comprises an amino acid substitution of the aspartic acid corresponding to position 242 of SEQ ID NO: 1. The TF antagonist also can be or comprise a human factor VII variant that comprises an amino acid substitution of the histidine corresponding to position 193 of SEQ ID NO: 1. A TF antagonist also can include any combination of these amino acid sequence variations. In particular aspects, the TF antagonist is or comprises a factor VII polypeptide selected from the list consisting of FVII-(K341A), FVII-(S344A), FVII-(D242A) and FVII-(H193A). The TF antagonist also can comprise,

consist essentially, or consist of a fragment of FVII or a fragment of a biologically active FVII variant.

The TF antagonist can also comprise any suitable combination of these and other TF antagonists. Thus, for example, the TF antagonist can comprise an antibody against TF and a 5 Factor VII variant.

The protein C polypeptide can be a wild-type protein C, such as human protein C, or a suitably active protein C-related polypeptide, such as a protein C polypeptide variant (i.e., a polypeptide differing from a wild-type protein C by one or more substitutions, deletions, and/or additions), or a derivative of protein C or protein C variant, or any suitable combination of such 10 polypeptides. The protein C polypeptide can be in a zymogen form or in activated form. In preferred embodiments, the protein C polypeptide is or comprises recombinant human protein C.

The invention described herein also provides a method of promoting, enhancing, and/or inducing one or more physiological responses in a host that are associated with the 15 treatment, amelioration, and/or prevention (e.g., reduction of the occurrence, onset, severity and/or promotion of recovery from) a thrombotic-related disease, coagulopathic related disease, respiratory disease, inflammatory disease, associated with TF, comprising administering a TF antagonist in combination with protein C or a protein C-related polypeptide to a host suffering from such a disease or at risk of developing such a disease or related disorder. The method thus 20 comprises the amelioration; reduction of onset, occurrence, severity, and/or spread of; prevention; and cure of such diseases. Such diseases and disorders can include deep venous thrombosis, arterial thrombosis, post surgical thrombosis, complications or disorders associated with the a coronary artery bypass graft (CABG) procedure, complications or disorders associated with percutaneous transdermal coronary angioplasty (PTCA), stroke, tumor metastasis, 25 inflammation, septic shock, hypotension, acute lung injury (ALI), Acute Respiratory Distress Syndrome (ARDS), pulmonary embolism, disseminated intravascular coagulation (DIC), sepsis, systemic inflammatory response syndrome (SIRS), vascular restenosis, platelet deposition, myocardial infarction, and angiogenesis-related disorders. The inventive method can also be used in the treatment or prophylactic treatment of mammals with atherosclerotic vessels at risk 30 for thrombosis; asthma, bronchitis, idiopathic pulmonary fibrosis, pneumonia, pulmonary edema, pulmonary obstructive disease, endotoxin induced lung damage, non cell lung cancer; inflammatory bowel disease, pancreatitis, trauma-induced shock, bronchial asthma, allergic rhinitis, rheumatoid arthritis, cystic fibrosis, stroke, acute bronchitis, chronic bronchitis, acute bronchiolitis, chronic bronchiolitis, osteoarthritis, gout, spondylarthropathy, ankylosing 35 spondylitis, Reiter's syndrome, psoriatic arthropathy, enteropathic spondylitis, juvenile arthropathy or juvenile ankylosing spondylitis, reactive arthropathy, infectious or post-infectious arthritis, gonococcal arthritis, tuberculous arthritis, viral arthritis, fungal arthritis, syphilitic

arthritis, Lyme disease, arthritis associated with vasculitic syndromes, polyarteritis nodosa, hypersensitivity vasculitis, Luegenec's granulomatosis, polymyalgia rheumatica, joint cell arteritis, calcium crystal deposition arthropathitis, pseudo gout, non-articular rheumatism, bursitis, tenosynovitis, epicondylitis (tennis elbow), carpal tunnel syndrome, repetitive use injury (e.g., 5 from typing), miscellaneous forms of arthritis, neuropathic joint disease (charco and joint), hemarthrosis (hemarthrosis), Henoch-Schonlein Purpura, hypertrophic osteoarthropathy, multicentric reticulohistiocytosis, arthritis associated with certain diseases, surcoilosis, hemochromatosis, sickle cell disease and other hemoglobinopathries, hyperlipoproteinemia, hypogammaglobulinemia, hyperparathyroidism, acromegaly, familial Mediterranean fever, 10 Behat's Disease, systemic lupus erythrematosis, relapsing, and multiple organ failure resulting from any of the preceding pathologic processes.

In one embodiment, the diseases or disorders treated by the inventive method are Respiratory disease and inflammatory disease. Respiratory and inflammatory diseases include lower respiratory diseases such as systemic inflammatory response syndrome, asthma, bronchitis, 15 acute lung injury, acute respiratory distress syndrome, idiopathic pulmonary fibrosis, pneumonia, pulmonary edema, pulmonary obstructive disease, endotoxin-induced lung damage, non cell lung cancer; inflammatory bowel disease, sepsis, septic shock, acute respiratory distress syndrome, pancreatitis, trauma-induced shock, bronchial asthma, allergic rhinitis, rheumatoid arthritis, cystic fibrosis, stroke, acute bronchitis, chronic bronchitis, acute bronchiolitis, chronic 20 bronchiolitis, osteoarthritis, gout, spondylarthropathitis, ankylosing spondylitis, Reiter's syndrome, psoriatic arthropathy, enteropathic spondylitis, juvenile arthropathy or juvenile ankylosing spondylitis, reactive arthropathy, infectious or post-infectious arthritis, gonococcal arthritis, tuberculous arthritis, viral arthritis, fungal arthritis, syphilitic arthritis, Lyme disease, 25 arthritis associated with "vasculitic syndromes," polyarteritis nodosa, hypersensitivity vasculitis, Luegenec's granulomatosis, polymyalgia rheumatica, joint cell arteritis, calcium crystal deposition arthropathitis, pseudo gout, non-articular rheumatism, bursitis, tenosynovitis, epicondylitis (tennis elbow), carpal tunnel syndrome, repetitive use injury (typing), miscellaneous forms of 30 arthritis, neuropathic joint disease (charco and joint), hemarthrosis (hemarthrosis), Henoch-Schonlein Purpura, hypertrophic osteoarthropathy, multicentric reticulohistiocytosis, arthritis associated with certain diseases, surcoilosis, hemochromatosis, sickle cell disease and other hemoglobinopathries, hyperlipoproteinemia, hypogammaglobulinemia, hyperparathyroidism, acromegaly, familial Mediterranean fever, Behat's Disease, systemic lupus erythrematosis, relapsing, and multiple organ failure resulting from any of the preceding pathologic processes.

Thrombotic or coagulopathic related diseases or disorders that may be treatable by the 35 inventive method include vascular diseases and inflammatory responses such as deep venous thrombosis, arterial thrombosis, post surgical thrombosis, complications or disorders associated with application of a coronary artery bypass graft (CABG), complications or disorders associated

with percutaneous transdermal coronary angioplasty (PTCA), stroke, tumor metastasis, inflammation, septic shock, hypotension, acute lung injury (ALI), Acute Respiratory Distress Syndrome (ARDS), pulmonary embolism, disseminated intravascular coagulation (DIC), sepsis, systemic inflammatory response syndrome (SIRS), vascular restenosis, platelet deposition, 5 myocardial infarction, angiogenesis-related disorders, or the treatment of mammals with atherosclerotic vessels at risk for thrombosis, and multiple organ failure resulting from any of the preceding pathologic processes.

In a preferred embodiment, the disease or disorder is one or more of systemic inflammatory response syndrome, acute lung injury, acute respiratory distress syndrome, 10 disseminated intravascular coagulation, sepsis, or multiple organ failure in association with any of the preceding syndromes.

The invention also provides a method of inhibiting of blood coagulation, thrombosis, pulmonary embolism, stroke, disseminated intravascular coagulation (DIC), platelet deposition, fibrin deposition in lungs and kidneys (e.g., deposition associated with gram-negative 15 endotoxemia), myocardial infarction, and inflammatory responses including acute lung injury, acute respiratory distress syndrome, and systemic inflammatory response syndrome comprising administering an effective amount of a TF antagonist and a protein C or protein C-related polypeptide or composition of the invention to a patient in need of the treatment of such a disease or disorder or at risk of developing such a disease or disorder.

20 The inventive method can be practiced with any suitable composition or by any suitable administration strategy described herein. Thus, in one aspect, the method comprises administering a TF antagonist and a protein C or protein C-related polypeptide in a single dose, whereas in other aspects the methods can be carried out by sequentially administered or simultaneously co-administration of the TF antagonist and protein C or protein C-related 25 polypeptide.

Any effective amount of TF antagonist and protein C or protein C-related polypeptide can be used in the inventive method. The particular dosage of these elements can vary with the condition to be treated, particular compositions used, and patient's physical characteristics. In one aspect of the invention, the TF antagonist and the protein C or protein C-related 30 polypeptide in the composition and/or used in the inventive method are present in a ratio by mass of between about 100:1 and about 1:100 (w/w TF antagonist: protein C). The method can be advantageously carried out by administering a pharmaceutical composition that is formulated for intravenous administration, preferably injection or infusion, and most preferably for injection.

35 The above-described elements and preferred aspects and features thereof will be described in further detail in the remaining sections of the detailed description of the invention.

Tissue factor antagonists

5 The terms "TF antagonist" or "TF antagonists", refer to any compound binding directly to TF and inhibiting the activation, or conversion, of factor X to factor Xa. Unless otherwise stated, in practicing the methods of present invention any such compound binding directly to tissue factor and inhibiting conversion of factor X to factor Xa may be used. This includes, without limitation, factor VII polypeptides having substantially reduced catalytic activity, inhibitory antibodies against TF, and fragments thereof. Desirably, the TF antagonists bind to tissue factor with high affinity and specificity but do not initiate blood coagulation.

10 In one embodiment of the invention, TF antagonists encompass those antagonists that exhibit at least about 10%, at least about 20%, at least about 25%, at least about 30%, at least about 40%, at least about 50%, at least about 60%, at least about 70%, at least about 75%, at least about 80%, at least about 90%, at least about 100%, at least about 110%, at least about 120%, or at least about 130%, of the specific TF-binding affinity of wild-type factor VIIa, when 15 tested in one or more of the TF binding assays as described in the present specification. In a preferred embodiment, the TF antagonists exhibit at least about 75% of the binding affinity of wild-type factor VIIa. The term "TF binding activity" as used herein means the ability of a VIIa polypeptide or TF antagonist to inhibit the binding of recombinant human  $^{125}\text{I}$ -VIIa to cell surface human TF. The TF binding activity may be measured as described in Assay 3 (of the 20 present specification).

25 In another embodiment, TF antagonists encompass those that exhibit less than about 50%, preferably less than about 25%, more preferably less than about 10%, or 5%, or 3%, or 2%, and most preferably less than about 1% of the specific activity of wild-type factor VIIa, when tested in one or more of a clotting assay, or proteolysis assay as described in the present specification.

30 The TF antagonists for use in the present invention include, without limitation, immunoglobulin molecules and fragments thereof that have the ability to specifically bind to an antigen (i.e., TF) such as (i) a Fab fragment, a monovalent fragment consisting of the VL, VH, CL and CH1 domains; (ii) F(ab)2 and F(ab')2 fragments; (iii) a Fd fragment consisting of the VH and CH1 domains; (iv) a Fv fragment consisting of the VL and VH domains of a single arm of an antibody, (v) a dAb fragment; (vi) an isolated complementarity determining region (CDR); (vii) single chain Fv (scFv); and (viii) diabodies. Included are also antibodies having variable and constant regions derived from human germ line immunoglobulin sequences; human antibodies including amino acid residues not encoded by human germ line immunoglobulin sequences (e.g., 35 mutations introduced by random or site-specific mutagenesis in vitro or by somatic mutation in vivo), for example in the CDRs and in particular CDR3; antibodies in which CDR sequences derived from the germ line of another mammalian species, such as a mouse, have been grafted

onto human framework sequences, e.g. the so-called humanized antibodies or human/mouse chimera antibodies.

The TF antagonists for use in the present invention also encompass, without limitation, factor VII polypeptides that has substantially reduced ability to catalyze the conversion of factor 5 X to factor Xa ("inactivated" factor VII polypeptides).

The terms "FVII polypeptide" or "FVII polypeptides" as used herein include, without limitation, native Factor VII, as well as factor VII-related polypeptides that have either been chemically modified relative to human factor VII and/or contain one or more amino acid sequence alterations relative to native Factor VII (i.e., Factor VII variants), and/or contain 10 truncated amino acid sequences relative to native Factor VII (i.e., Factor VII fragments).

It further encompasses polypeptides with a slightly modified amino acid sequence, for instance, polypeptides having a modified N-terminal end including N-terminal amino acid deletions or additions, and/or polypeptides that have been chemically modified relative to human factor VIIa. Such factor VII-related polypeptides may exhibit different properties relative 15 to native Factor VII, including stability, phospholipid binding, altered specific proteolytic activity, and the like. Factor VII-related polypeptides also include proteolytically inactive variants of Factor VII.

The terms "variant" or "variants", as used herein, is intended to designate human Factor VII having the sequence of SEQ ID NO: 1, wherein one or more amino acids of the parent 20 protein have been substituted by another amino acid and/or wherein one or more amino acids of the parent protein have been deleted and/or wherein one or more amino acids have been inserted in the protein and/or wherein one or more amino acids have been added to the parent protein.

The terms "Factor VII" or "FVII" are intended to include Factor VII polypeptides in their 25 uncleaved (zymogen) form as well as those that have been proteolytically processed to yield their respective bioactive forms, which may be designated "factor VIIa polypeptides" or "activated factor VII polypeptides"). Typically, FVII is cleaved between residues 152 and 153 to yield VIIa. The term "factor VII polypeptide" is also intended to encompass, without limitation, polypeptides having the amino acid sequence 1-406 of wild-type human Factor VII (as disclosed 30 in U.S. Patent No. 4,784,950), as well as wild-type Factor VII derived from other species, such as, e.g., bovine, porcine, canine, murine, and salmon Factor VII. It further encompasses natural allelic variations of Factor VII that may exist and occur from one individual to another. Also, degree and location of glycosylation or other post-translation modifications may vary depending on the chosen host cells and the nature of the host cellular environment. In one series of 35 embodiments, factor VII polypeptides include polypeptides that exhibit at least about 70 %, preferably at least about 80 %, more preferably at least about 90 %, and most preferable at

least about 95 %, of identity with the sequence of wild-type factor VII as disclosed in U.S. Patent No. 4,784,950.

Non-limiting examples of factor VII variants having substantially reduced or modified biological activity relative to wild-type factor VII include R152E-FVIIa (Wildgoose et al., Biochem 5 29:3413-3420, 1990), S344A-FVIIa (Kazama et al., J. Biol. Chem. 270:66-72, 1995), FFR-FVIIa (Holst et al., Eur. J. Vasc. Endovasc. Surg. 15:515-520, 1998), and factor VIIa lacking the Gla domain, (Nicolaisen et al., FEBS Letts. 317:245-249, 1993). Non-limiting examples also include human FVIIa, which has an amino acid substitution of the lysine corresponding to position 341 of SEQ ID NO: 1; human FVIIa, which has an amino acid substitution of the serine corresponding to position 10 10 344 of SEQ ID NO: 1; human FVIIa, which has an amino acid substitution of the aspartic acid corresponding to position 242 of SEQ ID NO: 1; human FVIIa, which has an amino acid substitution of the histidine corresponding to position 193 of SEQ ID NO: 1; FVII-(K341A); FVII-(S344A); FVII-(D242A); and FVII-(H193A). Non-limiting examples of chemically modified factor VII polypeptides and sequence variants are described, e.g., in U.S. Patent No. 5,997,864.

15 Non-limiting examples of FVII-derived peptides having TF/FVIIa antagonist activity are described in International patent applications WO 90/03390, WO 95/00541, WO 96/18653, and European Patent EP 500800.

The catalytic activity of Factor VIIa can be inhibited by chemical derivatization of the catalytic center, or triad. Derivatization may be accomplished by reacting Factor VII with an 20 irreversible inhibitor such as an organophosphor compound, a sulfonyl fluoride, a peptide halomethyl ketone or an azapeptide, or by acylation, for example, peptide chloromethylketones or peptidyl chloromethanes; azapeptides; acylating agents such as various guanidinobenzoate derivatives and 3-alkoxy-4-chloroisocoumarins; sulphonyl fluorides such as phenylmethylsulphonylfluoride (PMSF); diisopropylfluorophosphate (DFP); 25 tosylpropylchloromethyl ketone (TPCK); tosylsylchloromethyl ketone (TLCK); nitrophenylsulphonates; heterocyclic protease inhibitors such as isocoumarines, and coumarins.

Preferred peptide halomethyl ketones include Phe-Phe-Arg chloromethyl ketone, Phe-Phe-Arg chloromethylketone, D-Phe-Phe-Arg chloromethyl ketone, D-Phe-Phe-Arg chloromethylketone Phe-Pro-Arg chloromethylketone, D-Phe-Pro-Arg chloromethylketone, Phe-30 Pro-Arg chloromethylketone, D-Phe-Pro-Arg chloromethylketone, L-Glu-Gly-Arg chloromethylketone and D-Glu-Gly-Arg chloromethylketone, Dansyl-Phe-Phe-Arg chloromethyl ketone, Dansyl-Phe-Phe-Arg chloromethylketone, Dansyl-D-Phe-Phe-Arg chloromethyl ketone, Dansyl-D-Phe-Pro-Arg chloromethylketone, Dansyl-D-Phe-Arg chloromethylketone, Dansyl-Phe-Pro-Arg chloromethylketone, Dansyl-D-Phe-Pro-35 Arg chloromethylketone, Dansyl-L-Glu-Gly-Arg chloromethylketone and Dansyl-D-Glu-Gly-Arg chloromethylketone.

In preferred embodiments amino acid substitutions are made in the amino acid sequence of the Factor VII catalytic triad, defined herein as the regions which contain the amino acids which contribute to the Factor VIIa catalytic site. The substitutions, insertions or deletions in the catalytic triad are generally at or adjacent to the amino acids which form the catalytic site.

5 In the human and bovine Factor VII proteins, the amino acids which form a catalytic "triad" are Ser344, Asp242, and His193 (subscript numbering indicating position in SEQ ID NO:1). The catalytic sites in Factor VII from other mammalian species may be determined using presently available techniques including, among others, protein isolation and amino acid sequence analysis. Catalytic sites may also be determined by aligning a sequence with the sequence of  
10 other serine proteases, particularly chymotrypsin, whose active site has been previously determined (Sigler et al., J. Mol. Biol., 35:143-164 (1968), incorporated herein by reference), and therefrom determining from said alignment the analogous active site residues.

The amino acid substitutions, insertions or deletions are made so as to prevent or otherwise inhibit activation by the Factor VIIa of Factors X and/or IX. The Factor VII so modified  
15 should, however, also retain the ability to compete with authentic Factor VII and/or Factor VIIa for binding to tissue factor in the coagulation cascade. Such competition may readily be determined by means of, e.g., a clotting assay as described herein, or a competition binding assay using, e.g., a cell line having cell-surface tissue factor, such as the human bladder carcinoma cell line J82 (Sakai et al. J. Biol. Chem. 264: 9980-9988 (1989)).

20 The amino acids which form the catalytic site in Factor VII, such as Ser344, Asp242, and His193 in human and bovine Factor VII, may either be substituted or deleted. It is preferred to change only a single amino acid, thus minimizing the likelihood of increasing the antigenicity of the molecule or inhibiting its ability to bind tissue factor, however two or more amino acid changes (substitutions, additions or deletions) may be made and combinations of substitution(s),  
25 addition(s) and deletion(s) may also be made. In a preferred embodiment for human and bovine Factor VII, Ser344 is preferably substituted with Ala, but Gly, Met, Thr, or other amino acids can be substituted. It is preferred to replace Asp with Glu and to replace His with Lys or Arg. In general, substitutions are chosen to disrupt the tertiary protein structure as little as possible. One may introduce residue alterations as described above in the catalytic site of appropriate  
30 Factor VII sequence of human, bovine or other species and test the resulting protein for a desired level of inhibition of catalytic activity and resulting anticoagulant activity as described herein.

In preferred embodiments of human and bovine Factor VII, the active site residue Ser344 is modified, replaced with Gly, Met, Thr, or more preferably, Ala. Such substitution could be made separately or in combination with substitution(s) at other sites in the catalytic triad,  
35 which includes His193 and Asp242.

Further examples of TF antagonists that can be used in accordance with the present invention include, but are not limited to:

- Humanized TF-MAb blocking Factor X binding to the TF/FVIIa complex, CNTO-859 from Centocor (subsidiary of J&J); CNTO-859 is a humanized version of the murine TF-MAb, 5G8, developed by Scripps investigated for treatment of sepsis.

5 - Murine TF-MAb, 5G8 from Scripps.

- Humanized IgG1kappa MAb that binds to TF, hOAT from Sunol, currently in preclinical studies for cancer.

- Chimeric TF-MAb blocking FX binding to TF:FVIIa, cH36 from Sunol, currently in phase I for ALI, ARDS sepsis and coronary artery disease.

- TF-MAb blocking FVIIa binding to TF, I43 from Sunol.

10 - Humanized IgG4kappa MAb that binds to TF, hFAT from Sunol, investigated for the treatment of acute cardiovascular indications.

- Humanized mouse monoclonal antibody to TF, preventing FX and FIX binding, hATR-5 from Chugai Pharmaceuticals.

- Humanized anti-TF antibody F(ab')2 fragment inhibiting FIX and FX binding to human

15 TF:FVIIa, D3H44 from Genentech (Emory Univ.).

- Recombinant humanized monoclonal antibody F(ab')2 fragment that inhibits the TF/FVIIa complex from activating FIX and FX, PRO387474 from Genentech.

- Dimer FFR-FVIIa and PEGylated FFR-FVIIa as disclosed in WO 02/02764 (University of Minnesota).

20 - Immunoconjugate proteins constructed as a dimer of two identical chains, each having an effector domain which is the Fc region of an IgG1 immunoglobulin conjugated to a targeting domain which is a mutant form of factor VII that binds to tissue factor but does not initiate blood coagulation, as disclosed in WO 01/02439 (Garen).

25 Protein C polypeptides:

In practicing the present invention, any protein C polypeptide may be used that is effective in preventing or treating bleeding. This includes protein C polypeptides derived from blood or plasma, or produced by recombinant means.

The present invention encompasses protein C polypeptides, such as, e.g., those having the amino acid sequence disclosed in Beckmann et al., (Nucleic Acids Research 13:5233 (1985) (wild-type human protein C); in European patent No. EP 191606 and US patents Nos. US 4775624, US 5151268, and US 5270040 (Eli Lilly); in European patent No. EP 215548 and US patents Nos. US 5073609, US 5302529, and US 5516650 (ZymoGenetics); and in US 5009889 (Oklahoma Medical Research Foundation).

35 In some embodiments, the protein C polypeptide is human activated protein C, as disclosed, e.g., in US 4,981,952. In one series of embodiments, protein C polypeptides include polypeptides that exhibit at least about 10%, preferably at least about 30%, more preferably at

least about 50%, and most preferably at least about 70%, of the specific biological activity of human activated protein C when tested in one or more assays for Protein C activity. Protein C activity can, for example, be tested by means of the "Protein C assay" disclosed in the present invention. In one series of embodiments, protein C polypeptides include polypeptides that 5 exhibit at least about 90%, preferably at least about 100%, preferably at least about 120%, more preferably at least about 140%, and most preferably at least about 160%, of the specific biological activity of human activated protein C. In one series of embodiments, protein C polypeptides include polypeptides that exhibit at least about 70 %, preferably at least about 80 %, more preferably at least about 90 %, and most preferable at least about 95 %, of identity 10 with the sequence of wild-type protein C as disclosed in Beckmann et al., (Nucleic Acids Research 13:5233 (1985)).

As used herein, "protein C polypeptide" encompasses, without limitation, protein C, as well as protein C-related polypeptides. The term "protein C" is intended to encompass, without limitation, polypeptides having the amino acid sequence of wild-type human protein C (as 15 disclosed *supra*), as well as wild-type protein C derived from other species, such as, e.g., bovine, porcine, canine; murine, rat and salmon protein C, said protein C derived from blood or plasma, or produced by recombinant means. It further encompasses natural allelic variations of protein C that may exist and occur from one individual to another. Also, degree and location of glycosylation or other post-translation modifications may vary depending on the chosen host 20 cells and the nature of the host cellular environment. The term "protein C" is also intended to encompass protein C polypeptides in their uncleaved (zymogen) form, as well as those that have been proteolytically processed to yield their respective bioactive forms, which may be designated aPC.

"Protein C-related polypeptides" include, without limitation, protein C polypeptides 25 that have either been chemically modified relative to human protein C and/or contain one or more amino acid sequence alterations relative to human protein C (i.e., protein C variants), and/or contain truncated amino acid sequences relative to human protein C (i.e., protein C fragments). Such protein C-related polypeptides may exhibit different properties relative to human protein C, including stability, phospholipid binding, altered specific activity, and the like. 30 The term "protein C-related polypeptides" are intended to encompass such polypeptides in their uncleaved (zymogen) form, as well as those that have been proteolytically processed to yield their respective bioactive forms, which may be designated "aPC-related polypeptides" or "activated protein C-related polypeptides"

As used herein, "protein C-related polypeptides" encompasses, without limitation, 35 polypeptides exhibiting substantially the same or improved biological activity relative to wild-type human protein C, as well as polypeptides in which the protein C biological activity has been substantially modified or reduced relative to the activity of wild-type human protein C. These

polypeptides include, without limitation, protein C or activated protein C that has been chemically modified and protein C variants into which specific amino acid sequence alterations have been introduced that modify or disrupt the bioactivity of the polypeptide.

It further encompasses polypeptides with a slightly modified amino acid sequence, for 5 instance, polypeptides having a modified N-terminal end including N-terminal amino acid deletions or additions, and/or polypeptides that have been chemically modified relative to human protein C.

10 Protein C-related polypeptides, including variants, whether exhibiting substantially the same or better bioactivity than wild-type protein C, or, alternatively, exhibiting substantially modified or reduced bioactivity relative to wild-type protein C, include, without limitation, polypeptides having an amino acid sequence that differs from the sequence of wild-type protein C by insertion, deletion, or substitution of one or more amino acids.

15 Protein C-related polypeptides, including variants, encompass those that exhibit at least about 10%, at least about 20%, at least about 25%, at least about 30%, at least about 40%, at least about 50%, at least about 60%, at least about 70%, at least about 75%, at least about 80%, at least about 90%, at least about 100%, at least about 110%, at least about 120%, or at least about 130%, of the specific activity of wild-type activated protein C that has been produced in the same cell type, when tested in the "*Protein C assay*" as described in the present specification.

20 Protein C-related polypeptides, including variants, having substantially the same or improved biological activity relative to wild-type activated protein C encompass those that exhibit at least about 25%, preferably at least about 50%, more preferably at least about 75%, more preferably at least about 100%, more preferably at least about 110%, more preferably at least about 120%, and most preferably at least about 130% of the specific activity of wild-type activated protein C that has been produced in the same cell type, when tested in the "*Protein C assay*" as described in the present specification.

25 Protein C-related polypeptides, including variants, having substantially reduced biological activity relative to wild-type activated protein C are those that exhibit less than about 25%, preferably less than about 10%, more preferably less than about 5% and most preferably less than about 1% of the specific activity of wild-type activated protein C that has been produced in the same cell type when tested in the "*Protein C assay*" as described in the present specification.

30 In some embodiments the protein C polypeptides are protein C-related polypeptides, in particular variants, wherein the ratio between the activity of said protein C polypeptide and the activity of native human activated protein C (wild-type aPC) is at least about 1.25 when tested in the "*protein C Assay*" as described in the present specification; in other embodiments, the ratio

is at least about 2.0; in further embodiments, the ratio is at least about 4.0, in further embodiments, the ratio is at least about 8.0.

In some embodiments, the protein C polypeptide is human protein C, as disclosed, e.g., in Beckmann et al. (see above) (wild-type protein C). In some embodiments, the protein C polypeptide is human activated protein C. In one series of embodiments, the protein C polypeptides are protein C-related polypeptides that exhibits at least about 10%, preferably at least about 30%, more preferably at least about 50%, and most preferably at least about 70%, of the specific biological activity of human activated protein C. In some embodiments, the protein C polypeptides have an amino acid sequence that differs from the sequence of wild-type protein C by insertion, deletion, or substitution of one or more amino acids.

Protein C is commercially available as Xigris® from Eli Lilly, Inc. (recombinantly produced, activated human protein C), or Ceprotin from Baxter (plasma-derived human protein C). Examples of protein C polypeptides that can be used in accordance with the present invention include, but are not limited to Xigris® (Eli Lilly) and Ceprotin (Baxter).

15

#### Definitions

In the present context the three-letter or one-letter indications of the amino acids have been used in their conventional meaning as indicated in table 1. Unless indicated explicitly, the amino acids mentioned herein are L-amino acids. It is to be understood, that the first letter in, 20 for example, K337 represent the amino acid naturally present at the indicated position wild-type factor VII, and that, for example, [K337A]-FVIIa designates the FVII-variant wherein the amino acid represented by the one-letter code K naturally present in the indicated position is replaced by the amino acid represented by the one-letter code A.

25

Table 1: Abbreviations for amino acids:

Amino acid	Three-letter code	One-letter code
Glycine	Gly	G
Proline	Pro	P
Alanine	Ala	A
Valine	Val	V
Leucine	Leu	L
Isoleucine	Ile	I
Methionine	Met	M
Cysteine	Cys	C
Phenylalanine	Phe	F
Tyrosine	Tyr	Y
Tryptophan	Trp	W
Histidine	His	H
Lysine	Lys	K
Arginine	Arg	R
Glutamine	Gln	Q
Asparagine	Asn	N
Glutamic Acid	Glu	E
Aspartic Acid	Asp	D

The terms "factor VIIa" and "FVIIa" may be used interchangeably.

The term "active site" and the like when used herein with reference to FVIIa refer to the catalytic and zymogen substrate binding site, including the "S1" site of FVIIa as that term is defined by Schecter, I. and Berger, A., (1967) *Biochem. Biophys. Res. Commun.* 7:157-162.

The term "TF-mediated coagulation activity" means coagulation initiated by TF through the formation of the TF/FVIIa complex and its activation of FIX and Factor X to FIXa and FXa, respectively. TF-mediated coagulation activity is measured in an FXa generation assay. The term "FXa generation assay" as used herein is intended to mean any assay where activation of FX is measured in a sample comprising TF, FVIIa, FX, calcium and phospholipids. Examples of FXa generation assays are described in assay 1 and assay 2 (of the present specification).

A TF/FVIIa mediated or associated process or event, or a process or event associated with TF-mediated coagulation activity, is any event, which requires the presence of TF/FVIIa.

Such processes or events include, but are not limited to, formation of fibrin which leads to thrombus formation; platelet deposition; proliferation of smooth muscle cells (SMCs) in the vessel wall, such as, for example, in intimal hyperplasia or restenosis, which is thought to result from a complex interaction of biological processes including platelet deposition and thrombus formation, release of chemotactic and mitogenic factors, and the migration and proliferation of vascular smooth muscle cells into the intima of an arterial segment; and deleterious events associated with post-ischemic reperfusion, such as, for example, in patients with acute myocardial infarction undergoing coronary thrombolysis.

The general mechanism of blood clot formation is reviewed by Ganong, in *Review of Medical Physiology*, 13th ed., Lange, Los Altos Calif., pp 411-414 (1987). Coagulation requires the confluence of two processes, the production of thrombin which induces platelet aggregation and the formation of fibrin which renders the platelet plug stable. The process comprises several stages each requiring the presence of discrete proenzymes and profactors. The process ends in fibrin crosslinking and thrombus formation. Fibrinogen is converted to fibrin by the action of thrombin. Thrombin, in turn, is formed by the proteolytic cleavage of prothrombin. This proteolysis is effected by factor Xa which binds to the surface of activated platelets and in the presence of FVa and calcium, cleaves prothrombin. TF/FVIIa is required for the proteolytic activation of factor X by the extrinsic pathway of coagulation. Therefore, a process mediated by or associated with TF/FVIIa, or a TF-mediated coagulation activity includes any step in the coagulation cascade from the formation of the TF/FVIIa complex to the formation of a fibrin platelet clot and which initially requires the presence of TF/FVIIa. For example, the TF/FVIIa complex initiates the extrinsic pathway by activation of factor X to factor Xa, FIX to FIXa, and additional FVII to FVIIa. TF/FVIIa mediated or associated process, or TF-mediated coagulation activity can be conveniently measured employing standard assays such as those described in Roy,

S., (1991) J. Biol. Chem. 266:4665-4668, and O'Brien, D. et al., (1988) J. Clin. Invest. 82:206-212 for the conversion of factor X to factor Xa in the presence of TF/FVIIa and other necessary reagents.

The term "TF related diseases or disorders" as used herein means any disease or disorder, where TF is involved. This includes, but are not limited to diseases or disorders related to TF-mediated coagulation activity, thrombotic or coagulopathic related diseases or disorders or diseases or disorders such as inflammatory responses and chronic thromboembolic diseases or disorders associated with fibrin formation, including vascular disorders such as deep venous thrombosis, arterial thrombosis, post surgical thrombosis, coronary artery bypass graft (CABG), percutaneous transdermal coronary angioplasty (PTCA), stroke, tumor metastasis, angiogenesis, thrombolysis, arteriosclerosis and restenosis following angioplasty, acute and chronic indications such as inflammation, septic shock, septicemia, hypotension, adult respiratory distress syndrome (ARDS), disseminated intravascular coagulopathy (DIC), pulmonary embolism, platelet deposition, myocardial infarction, or the prophylactic treatment of mammals with atherosclerotic vessels at risk for thrombosis, and other diseases. The TF related diseases or disorders are not limited to in vivo coagulopathic disorders such as those named above, but includes ex vivo TF/FVIIa related processes such as coagulation that may result from the extracorporeal circulation of blood, including blood removed in-line from a patient in such processes as dialysis procedures, blood filtration, or blood bypass during surgery.

It should be noted that peptides, proteins and amino acids as used herein can comprise or refer to "natural", i.e., naturally occurring amino acids as well as "nonclassical" D-amino acids including, but not limited to, the D-isomers of the common amino acids,  $\alpha$ -isobutyric acid, 4-aminobutyric acid, hydroxyproline, sarcosine, citrulline, cysteic acid, t-butylglycine, t-butylalanine, phenylglycine, cyclohexylalanine,  $\beta$ -alanine, designer amino acids such as  $\beta$ -methyl amino acids,  $\alpha$ -methyl amino acids, N $\alpha$ -methyl amino acids, and amino acid analogues in general. In addition, the amino acids can include Abu, 2-amino butyric acid; (-Abu, 4-aminobutyric acid; (-Ahx, 6-aminohexanoic acid; Aib, 2-amino-isobutyric acid; (-Ala, 3-aminopropionic acid; Orn, ornithine; Hyp, trans-hydroxyproline; Nle, norleucine; Nva, norvaline.

The three-letter indication "GLA" as used herein means 4-carboxyglutamic acid ((-carboxyglutamate).

The terms "human tissue factor" or "human TF" as used herein, refers to the full length polypeptide receptor comprising the amino acid sequence 1-263 of native human tissue factor.

The term "antibody", as used herein, is intended to refer to immunoglobulin molecules and fragments thereof, which have the ability to specifically bind to an antigen (e.g., human TF). Full-length antibodies comprise four polypeptide chains, two heavy (H) chains and two light (L) chains inter-connected by disulfide bonds. Each heavy chain is comprised of a heavy chain variable region (abbreviated herein as HCVR or VH) and a heavy chain constant region. The heavy chain constant region is comprised of three domains, CH1, CH2, and CH3. Each light chain

is comprised of a light chain variable region (abbreviated herein as LCVR or VL) and a light chain constant region. The light chain constant region is comprised of one domain, CL. The VH and VL regions can be further subdivided into regions of hypervariability, termed complementarity determining regions (CDR), interspersed with regions that are more conserved, termed framework regions (FR). Each VH and VL is composed of three CDRs and four FRs, arranged from amino-terminus to carboxy-terminus in the following order: FR1, CDR1, FR2, CDR2, FR3, CDR3, FR4. Thus, included within the definition of an antibody are also one or more fragments of an antibody that retain the ability to specifically bind to an antigen (e.g., human TF). It has been shown that the antigen-binding function of an antibody can be performed by fragments of a full-length antibody. Examples of binding fragments encompassed within the term "antibody" include (i) a Fab fragment, a monovalent fragment consisting of the VL, VH, CL and CH1 domains; (ii) F(ab)2 and F(ab')2 fragments, a bivalent fragment comprising two Fab fragments linked by a disulfide bridge at the hinge region; (iii) a Fd fragment consisting of the VH and CH1 domains; (iv) a Fv fragment consisting of the VL and VH domains of a single arm of an antibody, (v) a dAb fragment (Ward et al., (1989) *Nature* 341:544-546 ), which consists of a VH domain; and (vi) an isolated complementarity determining region (CDR). Furthermore, although the two domains of the Fv fragment, VL and VH, are coded for by separate genes, they can be joined, using recombinant methods, by a synthetic linker that enables them to be made as a single protein chain in which the VL and VH regions pair to form monovalent molecules (known as single chain Fv (scFv); see e.g., Bird et al. (1988) *Science* 242:423-426: and Huston et al. (1988) *Proc. Natl. Acad. Sci. USA* 85:5879-5883). Such single chain antibodies are also intended to be encompassed within the term "antibody". Other forms of single chain antibodies, such as diabodies are also encompassed. Diabodies are bivalent, bispecific antibodies in which VH and VL domains are expressed on a single polypeptide chain, but using a linker that is too short to allow for pairing between the two domains on the same chain, thereby forcing the domains to pair with complementary domains of another chain and creating two antigen binding sites (see e.g., Hollinger, P., et al. (1993) *Proc. Natl. Acad. Sci. USA* 90:6444-6448; Poljak, R. J., et al. (1994) *Structure* 2:1121-1123). It is understood that human TF may have one or more antigenic determinants comprising (1) peptide antigenic determinants which consist of single peptide chains within human TF, (2) conformational antigenic determinants which consist of more than one spatially contiguous peptide chains whose respective amino acid sequences are located disjointedly along the human TF polypeptide sequence; and (3) post-translational antigenic determinants which consist, either in whole or part, of molecular structures covalently attached to human TF after translation, such as carbohydrate groups, or the like.

The terms "human antibody", "human antibodies", "human TF antibody", and "human TF antibodies", as used herein, is intended to include antibodies having variable and constant regions derived from human germ line immunoglobulin sequences. The human antibodies of

the invention may include amino acid residues not encoded by human germ line immunoglobulin sequences (e.g., mutations introduced by random or site-specific mutagenesis in vitro or by somatic mutation in vivo), for example in the CDRs and in particular CDR3. The term "humanized antibody", as used herein, is intended to include antibodies in which CDR sequences 5 derived from the germ line of another mammalian species, such as a mouse, have been grafted onto human framework sequences; the term "humanized antibody" is exchangeable with "chimeric antibody", and "chimera antibodies", (e.g., human/mouse chimera antibodies").

By "catalytically inactivated in the active site of the FVIIa polypeptide" is meant that a factor VIIa inhibitor is bound to the factor VIIa polypeptide and decreases or prevents the factor 10 VIIa-catalyzed conversion of factor X to factor Xa. An FVIIa inhibitor may be identified as a substance, which reduces the amidolytic activity by at least 50% at a concentration of the substance at 400  $\mu$ M in the FVIIa amidolytic assay described by Persson et al. (Persson et al., J. Biol. Chem. 272: 19919-19924 (1997)). Preferred are substances reducing the amidolytic activity by at least 50% at a concentration of the substance at 300  $\mu$ M; more preferred are substances 15 reducing the amidolytic activity by at least 50% at a concentration of the substance at 200  $\mu$ M.

The "FVIIa inhibitor" may be selected from any one of several groups of FVIIa directed inhibitors. Such inhibitors are broadly categorized for the purpose of the present invention into i) inhibitors which reversibly bind to FVIIa and are cleavable by FVIIa, ii) inhibitors which reversibly bind to FVIIa but cannot be cleaved, and iii) inhibitors which irreversibly bind to FVIIa. 20 For a review of inhibitors of serine proteases see Proteinase Inhibitors (Research Monographs in cell and Tissue Physiology; v. 12) Elsevier Science Publishing Co., Inc., New York (1990).

The FVIIa inhibitor moiety may also be an irreversible FVIIa serine protease inhibitor. Such irreversible active site inhibitors generally form covalent bonds with the protease active site. Such irreversible inhibitors include, but are not limited to, general serine protease inhibitors 25 such as peptide chloromethylketones (see, Williams et al., J. Biol. Chem. 264:7536-7540 (1989)) or peptidyl chloromethanes; azapeptides; acylating agents such as various guanidinobenzoate derivatives and the 3-alkoxy-4-chloroisocoumarins; sulphonyl fluorides such as phenylmethylsulphonylfluoride (PMSF); diisopropylfluorophosphate (DFP); tosylpropylchloromethyl ketone (TPCK); tosyllysylchloromethyl ketone (TLCK); 30 nitrophenylsulphonates and related compounds; heterocyclic protease inhibitors such as isocoumarines, and coumarins.

Examples of peptidic irreversible FVIIa inhibitors include, but are not limited to, Phe-Phe-Arg chloromethyl ketone, Phe-Phe-Arg chloromethylketone, D-Phe-Phe-Arg chloromethyl ketone, D-Phe-Phe-Arg chloromethylketone Phe-Pro-Arg chloromethylketone, D- 35 Phe-Pro-Arg chloromethylketone, Phe-Pro-Arg chloromethylketone, D-Phe-Pro-Arg chloromethylketone, L-Glu-Gly-Arg chloromethylketone, and D-Glu-Gly-Arg chloromethylketone.

Examples of FVIIa inhibitors also include benzoxazinones or heterocyclic analogues thereof such as described in PCT/DK99/00138.

Examples of other FVIIa inhibitors include, but are not limited to, small peptides such as for example Phe-Phe-Arg, D-Phe-Phe-Arg, Phe-Phe-Arg, D-Phe-Phe-Arg, Phe-Pro-Arg, D-Phe-Pro-Arg, Phe-Pro-Arg, D-Phe-Pro-Arg, L- and D-Glu-Gly-Arg; peptidomimetics; benzamidine systems; heterocyclic structures substituted with one or more amidino groups; aromatic or heteroaromatic systems substituted with one or more C(=NH)NHR groups in which R is H, C1-3alkyl, OH or a group which is easily split of *in vivo*.

The terms "aPC", "APC", "Activated Protein C", "raPC", "rAPC", and "recombinant Activated Protein C" are synonymous for the purpose and practice of this invention and can be used interchangeably.

**Protein C Activity:** any property of activated human Protein C or its derivatives responsible for proteolytic, amidolytic, esterolytic, and biological (anticoagulant or profibrinolytic) activities. Methods for testing for Protein C anticoagulant and amidolytic activity are well known in the art, i.e., see Grinnell et. al., 1987, Bio/Technology 5:1189-1192.

**RhaPC:** Recombinant activated human protein C, produced by activating r-HPC in vitro or by direct secretion of the activated form of Protein C from prokaryotic cells, eukaryotic cells, or from transgenic animals.

**Zymogen:** an enzymatically inactive precursor of a proteolytic enzyme. Protein C zymogen, as used herein, refers to secreted, inactive forms, whether one chain or two chain, of protein C.

**Thrombotic or coagulopathic related diseases or disorders:** the term includes vascular diseases and inflammatory responses including, without limitation, deep venous thrombosis, arterial thrombosis, post surgical thrombosis, coronary artery bypass graft (CABG), percutaneous transdermal coronary angioplasty (PTCA), stroke, tumor metastasis, inflammation, septic shock, hypotension, acute lung injury (ALI), Acute Respiratory Distress Syndrome (ARDS), pulmonary embolism, disseminated intravascular coagulation (DIC), sepsis, systemic inflammatory response syndrome (SIRS), vascular restenosis, platelet deposition, myocardial infarction, angiogenesis, or the treatment of mammals with atherosclerotic vessels at risk for thrombosis, and multiple organ failure resulting from any of the preceding pathologic processes.

**Respiratory Diseases or disorders:** exemplified by lower respiratory diseases such as systemic inflammatory response syndrome, asthma, bronchitis, acute lung injury, acute respiratory distress syndrome, idiopathic pulmonary fibrosis, pneumonia, pulmonary edema, pulmonary obstructive disease, endotoxin induced lung damage, non cell lung cancer, and multiple organ failure resulting from any of the preceding pathologic processes.

**Inflammatory Diseases or disorders:** refers to diseases such as inflammatory bowel disease, sepsis, septic shock, acute respiratory distress syndrome, pancreatitis, trauma-induced

shock, bronchial asthma, allergic rhinitis, rheumatoid arthritis, cystic fibrosis, stroke, acute bronchitis, chronic bronchitis, acute bronchiolitis, chronic bronchiolitis, osteoarthritis, gout, spondylarthropathis, ankylosing spondylitis, Reiter's syndrome, psoriatic arthropathy, enteropathic spondylitis, juvenile arthropathy or juvenile ankylosing spondylitis, reactive 5 arthropathy, infectious or post-infectious arthritis, gonococcal arthritis, tuberculous arthritis, viral arthritis, fungal arthritis, syphilitic arthritis, Lyme disease, arthritis associated with "vasculitic syndromes," polyarteritis nodosa, hypersensitivity vasculitis, Luegenec's granulomatosis, polymyalgia rheumatica, joint cell arteritis, calcium crystal deposition arthropathis, pseudo gout, non-articular rheumatism, bursitis, tenosynovitis, epicondylitis (tennis elbow), carpal 10 tunnel syndrome, repetitive use injury (typing), miscellaneous forms of arthritis, neuropathic joint disease (charco and joint), hemarthrosis (hemarthrosis), Henoch-Schonlein Purpura, hypertrophic osteoarthropathy, multicentric reticulohistiocytosis, arthritis associated with certain diseases, surcoilosis, hemochromatosis, sickle cell disease and other hemoglobinopathies, hyperlipoproteinemia, hypogammaglobulinemia, hyperparathyroidism, acromegaly, familial 15 Mediterranean fever, Behat's Disease, systemic lupus erythrematosis, relapsing, and multiple organ failure resulting from any of the preceding pathologic processes.

The phrase "therapeutically effective interval" is a period of time beginning when one of either (a) the TF antagonist or (b) Protein C or protein C-related polypeptide is administered to a mammal and ending at the limit of the beneficial effect in preventing or ameliorating 20 respiratory or inflammatory disease or associated organ failure of (a) or (b).

"Sole" agents or factors as used herein refers to situations in which the TF antagonist and the protein C or protein C-related polypeptide, taken together, are the only haemostatic agents, or active haemostatic agents, or coagulation factors contained in the pharmaceutical composition or kit, or are the only haemostatic agents, or active haemostatic agents, or 25 coagulation factors administered to the patient in the course of a particular treatment, such as, e.g., in the course of a particular bleeding episode. It will be understood that these situations encompass those in which other haemostatic agents or coagulation factors, as applicable, are not present in either sufficient quantity or activity so as to significantly influence one or more coagulation parameters.

30 Clot lysis time, clot strength, fibrin clot formation, and clotting time are clinical parameters used for assaying the status of patient's haemostatic system. Blood samples are drawn from the patient at suitable intervals and one or more of the parameters are assayed by means of, e.g., thromboelastography as described by, e.g., Meh et al., Blood Coagulation & Fibrinolysis 2001;12:627-637; Vig et al., Hematology, Vol. 6 (3) pp. 205-213 (2001); Vig et al., Blood coagulation & fibrinolysis, 35 Vol. 12 (7) pp. 555-561 (2001) Oct; Glidden et al., Clinical and applied thrombosis/hemostasis, Vol. 6 (4) pp. 226-233 (2000) Oct; McKenzie et al., Cardiology, Vol. 92 (4) pp. 240-247 (1999) Apr; or Davis et al., Journal of the American Society of Nephrology, Vol. 6 (4) pp. 1250-1255 (1995).

In this context, the term "treatment" is meant to include both prevention of an expected unwanted clotting, and regulation of an already occurring clotting. Prophylactic administration of a preparation of a TF antagonist and a protein C polypeptide is thus included in the term "treatment".

5 The term "subject" as used herein is intended to mean any animal, in particular mammals, such as humans, and may, where appropriate, be used interchangeably with the term "patient".

The TF antagonist and protein C or protein C-related polypeptide as defined in the present specification may be administered simultaneously or sequentially. The factors may be supplied in single-dosage form wherein the single-dosage form contains both coagulation 10 factors, or in the form of a kit-of-parts comprising a preparation of a TF antagonist as a first unit dosage form and a preparation of protein C or protein C-related polypeptide as a second unit dosage form. Whenever a first or second or third, etc., unit dose is mentioned throughout this specification this does not indicate the preferred order of administration, but is merely done for convenience purposes

15 By "simultaneous" dosing of a preparation of a TF antagonist and a preparation of protein C or protein C-related polypeptide is meant administration of the coagulation factor proteins in single-dosage form, or administration of a first coagulation factor protein followed by administration of a second coagulation factor protein with a time separation of no more than 15 minutes, preferably 10, more preferred 5, more preferred 2 minutes. Either factor may be 20 administered first.

By "sequential" dosing is meant administration of a first coagulation factor protein followed by administration of a second coagulation factor protein with a time separation of up to 2 hours, preferably from 1 to 2 hours, more preferred up to 1 hour, more preferred from 30 25 minutes to 1 hour, more preferred up to 30 minutes, more preferred from 15 to 30 minutes. Either of the two unit dosage form, or coagulation factor proteins, may be administered first. Preferably, both products are injected through the same intravenous access.

30 By "APTT" or "aPTT" is meant the activated partial thromboplastin time (described by, e.g., Proctor RR, Rapaport SI: The partial thromboplastin time with kaolin; a simple screening test for first-stage plasma clotting factor deficiencies. Am J Clin Pathol 36:212, 1961).

"Half-life" refers to the time required for the plasma concentration of a TF antagonist or protein C or a protein C-related polypeptide to decrease from a particular value to half of that value.

35 The total amount of protein in a preparation may be measured by generally known methods, e.g., by measuring optical density. Amounts of protein C polypeptides or factor VII protein ("antigen") may be measured by generally known methods such as standard Elisa immunoassays. In general terms, such assay is conducted by contacting, e.g., a solution of the

protein C-containing preparation with an anti-protein C antibody immobilized onto the ELISA plate, subsequently contacting the immobilized antibody-protein C complex with a second anti protein C antibody carrying a marker, the amounts of which, in a third step, are measured. The amounts of each coagulation factor may be measured in a similar way using appropriate 5 antibodies. The total amount of coagulation factor protein present in a preparation is determined by adding the amounts of the individual coagulation factor proteins. In one embodiment, the preparation comprises isolated coagulation factor. In another embodiment the preparation is free of coagulation factor II and coagulation factor IIa (prothrombin and thrombin) and/or factor X or Xa.

10 As used herein, the term "isolated" refers to coagulation factors, e.g., protein C or protein C-related polypeptides that have been separated from the cell in which they were synthesized or the medium in which they are found in nature (e.g., plasma or blood). Separation of polypeptides from their cell of origin may be achieved by any method known in the art, including, without limitation, removal of cell culture medium containing the desired product 15 from an adherent cell culture; centrifugation or filtration to remove non-adherent cells; and the like. Separation of polypeptides from the medium in which they naturally occur may be achieved by any method known in the art, including, without limitation, affinity chromatography, such as, e.g., on an anti-factor VII or anti-protein C antibody column, respectively; hydrophobic interaction chromatography; ion-exchange chromatography; size 20 exclusion chromatography; electrophoretic procedures (e.g., preparative isoelectric focusing (IEF)), differential solubility (e.g., ammonium sulfate precipitation), or extraction and the like.

Within the present invention an "effective amount" of a TF antagonist and a protein C or protein C-related polypeptide is defined as the amount of a TF antagonist, e.g., inactivated FVIIa or a TF antibody, and a protein C polypeptide that together suffices to prevent or reduce 25 bleeding or blood loss, so as to cure, alleviate or partially arrest the disease and its complications.

The phrase "therapeutically effective combination," used in the practice of this invention, means administration of both (a) a TF antagonist and (b) protein C or a protein C-related polypeptide, either simultaneously or separately.

30 The term, "Active Ingredient" as used herein refers to a combination of (a) a TF antagonist and (b) Protein C or a protein C-related polypeptide co-present in a pharmaceutical formulation for the delivery of a treatment regimen that applies this invention.

The term, "injectable liquid carrier" refers to a liquid medium containing either or both of (a) a TF antagonist, or (b) Protein C or a protein C-related polypeptide; wherein (a) and (b) are independently dissolved, suspended, dispersed, or emulsified in the liquid medium.

35

Abbreviations

TF tissue factor

FVII	factor VII in its single-chain, unactivated form
FVIIa	factor VII in its activated form
rFVIIa	recombinant factor VII in its activated form
APC	Activated human Protein C, also called, Activated Protein C.
5 HPC	human Protein C zymogen.
rhPC	recombinant human Protein C zymogen.
APTT	activated partial thromboplastin time.

10 *Preparation of compounds:*

Methods for preparing recombinant proteins including conventional molecular biology, microbiology, and recombinant DNA techniques are within the skill of the art. Such techniques are explained fully in the literature. See, e.g., Sambrook, Fritsch & Maniatis, *Molecular Cloning: A Laboratory Manual*, Second Edition (1989) Cold Spring Harbor Laboratory Press, Cold Spring Harbor, New York (herein "Sambrook et al., 1989") *DNA Cloning: A Practical Approach*, Volumes I and II /D.N. Glover ed. 1985); *Oligonucleotide Synthesis* (M.J. Gait ed. 1984); *Nucleic Acid Hybridization* (B.D. Hames & S.J. Higgins eds (1985)); *Transcription And Translation* (B.D. Hames & S.J. Higgins, eds. (1984)); *Animal Cell Culture* (R.I. Freshney, ed. (1986)); *Immobilized Cells And Enzymes* (IRL Press, (1986)); B. Perbal, *A Practical Guide To Molecular Cloning* (1984).

20 Briefly, DNA sequences encoding a specific protein (e.g., protein C or human FVII) may be isolated by preparing a genomic or cDNA library and screening for DNA sequences coding for all or part of the protein by hybridization using synthetic oligonucleotide probes in accordance with standard techniques (cf. Sambrook et al., above). For the present purpose, the DNA sequence encoding the protein is preferably of human origin, i.e. derived from a human genomic DNA or cDNA library.

25 Polypeptide variants may be made by amino acid sequence alterations of the polypeptide, which may be accomplished by a variety of techniques. Modification of the DNA sequence may be by site-specific mutagenesis. Techniques for site-specific mutagenesis are well known in the art and are described by, for example, Zoller and Smith (DNA 3:479-488, 1984).

30 The DNA sequences encoding the polypeptide may also be prepared synthetically by established standard methods, e.g. the phosphoamidite method described by Beaucage and Caruthers, *Tetrahedron Letters* 22 (1981), 1859 - 1869, or the method described by Matthes et al., *EMBO Journal* 3 (1984), 801 - 805. According to the phosphoamidite method, oligonucleotides are synthesized, e.g. in an automatic DNA synthesizer, purified, annealed, ligated and cloned in 35 suitable vectors.

The DNA sequences may also be prepared by polymerase chain reaction using specific primers, for instance as described in US 4,683,202, Saiki et al., *Science* 239 (1988), 487 - 491, or Sambrook et al., *supra*.

The host cell into which the DNA sequences encoding the polypeptides is introduced 5 may be any cell, which is capable of producing the posttranslational modified human FVII polypeptides and includes yeast, fungi and higher eukaryotic cells. Methods of transfecting mammalian cells and expressing DNA sequences introduced in the cells are described in e.g. Kaufman and Sharp, *J. Mol. Biol.* 159 (1982), 601 - 621; Southern and Berg, *J. Mol. Appl. Genet.* 1 (1982), 327 - 341; Loyter et al., *Proc. Natl. Acad. Sci. USA* 79 (1982), 422 - 426; Wigler et al., *Cell* 14 10 (1978), 725; Corsaro and Pearson, *Somatic Cell Genetics* 7 (1981), 603, Graham and van der Eb, *Virology* 52 (1973), 456; and Neumann et al., *EMBO J.* 1 (1982), 841 - 845.

Methods for producing antibodies and fragments of antibodies are generally known in the art, see, e.g., Harboe and Ingild, In N.H. Axelsen, J. Krøll, and B. Weeks, editors, *A Manual of Quantitative Immunoelectrophoresis*, Blackwell Scientific Publications, 1973, Chapter 23, or 15 Johnstone and Thorpe, *Immunochemistry in Practice*, Blackwell Scientific Publications, 1982 (more specifically pages 27-31). Preferably, the antibodies are monoclonal antibodies. Monoclonal antibodies may be prepared, e.g., according to the methods of E. Harlow and D. Lane, editors, 1988, *Antibodies, A Laboratory Manual*, Cold Spring Harbor Press, Cold Spring Harbor, New York.

20 Methods of preparing human antibodies against human TF are, for example, described in International application No. PCT/DK02/00644

#### **Factor VII polypeptides**

For the preparation of recombinant human FVII polypeptides, a cloned wild-type FVII 25 DNA sequence is used. This sequence may be modified to encode a desired FVII variant. The complete nucleotide and amino acid sequences for human FVII are known; see U.S. Pat. No. 4,784,950, where the cloning and expression of recombinant human FVII is described. The bovine FVII sequence is described in Takeya et al., *J. Biol. Chem.* 263:14868-14872 (1988), which is incorporated by reference herein.

30 DNA sequences for use will typically encode a pre-pro peptide at the amino-terminus of the FVII protein to obtain proper post-translational processing (e.g. gamma-carboxylation of glutamic acid residues) and secretion from the host cell. The pre-pro peptide may be that of FVII or another vitamin K-dependent plasma protein, such as factor IX, factor X, prothrombin, protein C or protein S. As will be appreciated by those skilled in the art, additional modifications 35 can be made in the amino acid sequence of FVII where those modifications do not significantly have impact on the ability of the protein to act as a coagulation factor. For example, FVII modified in the catalytic triad can also be modified in the activation cleavage site to inhibit the

conversion of zymogen FVII into its activated two-chain form, as generally described in U.S. Pat. No. 5,288,629.

Factor VII polypeptides for use in the present invention may be prepared, e.g., as described in International Applications Nos. WO 92/15686, WO 94/27631 and WO 96/12800; 5 Wildgoose et al., Biochem 29:3413-3420, 1990; Kazama et al., J. Biol. Chem. 270:66-72, 1995; Holst et al., Eur. J. Vasc. Endovasc. Surg. 15:515-520, 1998; and Nicolaisen et al., FEBS Letts. 317:245-249, 1993.

FVII polypeptides produced as described above may be purified by affinity chromatography on an anti-FVII antibody column. The immunoabsorption column comprise a 10 high-specificity monoclonal antibody, such as, e.g., a calcium-dependent monoclonal antibody as described by Wakabayashi et al., J. Biol. Chem. 261:11097-11108, (1986) and Thim et al., Biochem. 27: 7785-7793, (1988). Additional purification may be achieved by conventional chemical 15 purification means, such as high performance liquid chromatography. Other methods of purification, including barium citrate precipitation, are known in the art, and may be applied to the purification of the FVIIa described herein (see, generally, Scopes, R., *Protein Purification*, Springer-Verlag, N.Y., 1982).

Conversion of single-chain FVII to active two-chain FVIIa may be achieved using factor Xlla as described by Hedner and Kisiel (1983, J. Clin. Invest. 71: 1836-1841), or using other 20 proteases having trypsin-like specificity (Kisiel and Fujikawa, Behring Inst. Mitt. 73: 29-42, 1983). Alternatively FVII may be autoactivated by passing it through an ion-exchange chromatography column, such as mono Q.RTM. (Pharmacia Fine Chemicals) or the like (Bjoern et al., 1986, Research Disclosures 269:564-565).

*Preparation of the Activated Protein C Ingredient.*

25 The cloning of human protein C has been described by Beckmann et al., (Nucleic Acids Research 13:5233 (1985) (wild-type human protein C). The expression of recombinant human Protein C in human kidney 293 cells has been described by Grinnell et al. (BioTechnology 5:1189-1192 (1987)). Recombinant human Protein C (r-hPC) may be produced by techniques well known to the skilled artisan such as those set forth in Yan, U.S. Patent No. 4,981,952. The gene 30 encoding human Protein C is disclosed and claimed in Bang et al., U.S. Patent No. 4,775,624. A plasmid useful to express human Protein C in 293 cells (pLPC) is disclosed in Bang et al., U.S. Patent No. 4,992,373; the construction of plasmid pLPC is also described in European Patent Publication No. 0 445 939 and in Grinnell et al., 1987, Bio/Technology 5:1189-1192. Briefly, the plasmid was transfected into 293 cells; stable transformants were identified, subcultured and 35 grown in serum-free media. After fermentation, cell-free medium was obtained by microfiltration.

Production of recombinant protein C is also described in European patent No. EP 191606 and US patents Nos. US 4775624, US 5151268, and US 5270040 (Eli Lilly); in European patent No. EP 215548 and US patents Nos. US 5073609, US 5302529, and US 5516650 (ZymoGenetics); and in US 5009889 (Oklahoma Medical Research Foundation).

5 Production and isolation of Protein C is also described by Haley et al., *J. Biol. Chem.*, 264; 16303, 1989, and Turkay et al., *Thromb. Haemost.* 81; 727 1999.

#### *Pharmaceutical Compositions and Methods of Use*

10 The preparations of the present invention may be used to treat thrombotic and coagulopathic related diseases or disorders, respiratory diseases or disorders, and inflammatory diseases or disorders including, without limitation, deep venous thrombosis, arterial thrombosis, post surgical thrombosis, coronary artery bypass graft (CABG), percutaneous transdermal coronary angioplasty (PTCA), stroke, tumor metastasis, inflammation, septic shock, hypotension, 15 acute lung injury (ALI), Acute Respiratory Distress Syndrome (ARDS), pulmonary embolism, disseminated intravascular coagulation (DIC), sepsis, systemic inflammatory response syndrome (SIRS), vascular restenosis, platelet deposition, myocardial infarction, angiogenesis, or the prophylactic treatment of mammals with atherosclerotic vessels at risk for thrombosis; asthma, bronchitis, idiopathic pulmonary fibrosis, pneumonia, pulmonary edema, pulmonary obstructive 20 disease, endotoxin induced lung damage, non cell lung cancer; inflammatory bowel disease, pancreatitis, trauma-induced shock, bronchial asthma, allergic rhinitis, rheumatoid arthritis, cystic fibrosis, stroke, acute bronchitis, chronic bronchitis, acute bronchiolitis, chronic bronchiolitis, osteoarthritis, gout, spondylarthropathy, ankylosing spondylitis, Reiter's syndrome, psoriatic arthropathy, enteropathic spondylitis, juvenile arthropathy or juvenile 25 ankylosing spondylitis, reactive arthropathy, infectious or post-infectious arthritis, gonococcal arthritis, tuberculous arthritis, viral arthritis, fungal arthritis, syphilitic arthritis, Lyme disease, arthritis associated with "vasculitic syndromes," polyarteritis nodosa, hypersensitivity vasculitis, Luegenec's granulomatosis, polymyalgia rheumatica, joint cell arteritis, calcium crystal deposition arthropathy, pseudo gout, non-articular rheumatism, bursitis, tenosynovitis, epicondylitis 30 (tennis elbow), carpal tunnel syndrome, repetitive use injury (typing), miscellaneous forms of arthritis, neuropathic joint disease (charcot and joint), hemarthrosis (hemarthrosis), Henoch-Schonlein Purpura, hypertrophic osteoarthropathy, multicentric reticulohistiocytosis, arthritis associated with certain diseases, surcoiosis, hemochromatosis, sickle cell disease and other hemoglobinopathies, hyperlipoproteinemia, hypogammaglobulinemia, hyperparathyroidism, 35 acromegaly, familial Mediterranean fever, Behcet's Disease, systemic lupus erythematosus, relapsing, and multiple organ failure resulting from any of the preceding pathologic processes.

The essential ingredients (a) a TF antagonist and (b) Protein C or a protein C-related polypeptide are present in the formulation in such proportion that a dose of the formulation provides an amount of each ingredient that together is a pharmaceutically effective amount to the patient being treated. The dose of composition of the invention to be administered is 5 determined depending upon age, body weight, symptom, the desired therapeutic effect, the route of administration, and the duration of the treatment etc. Typically, the weight ratio of TF antagonist and the amount of protein C or protein C-related polypeptide (e.g., APC or a biologically active fragment or variant thereof) may vary from a ratio of between about 1:100 to about 100:1 (w/w). The ratio of TF antagonist to protein C or protein C-related polypeptide (e.g., 10 APC or biologically active fragment or variant) may thus be, e.g., about 1:100, or 1:90, or 1:80, or 1:70 or 1:60, or 1:50, or 1:40, or 1:30, or 1:20, or 1:10, or 1:5, or 1:2, or 1:1, or 2:1, or 5:1, or 10:1, or 20:1, or 30:1, or 40:1, or 50:1, or 60:1, or 70:1, or 80:1, or 90:1, or 100:1; or between about 1:90 to about 1:1, or between about 1:80 to about 1:2, or between about 1:70 to about 1:5, or between about 1:60 to about 1:10, or between about 1:50 to about 1:25, or between about 1:40 15 to about 1:30, or between about 90:1 to about 1:1, or between about 80:1 to about 2:1, or between about 70:1 to about 5:1, or between about 60:1 to about 10:1, or between about 50:1 to about 25:1, or between about 40:1 to about 30:1; or between about 10:1 to about 1:10, or between about 5:1 to about 1:5.

The dose of the TF antagonist ranges from about 0.05 mg to about 500 mg/day, e.g., 20 from about 1 mg to about 200 mg/day, or, e.g., from about 5 mg to about 175 mg/day for a 70-kg subject as loading and maintenance doses, depending on the weight of the subject, the condition and the severity of the condition.

The dose of the protein C or protein C-related polypeptide ranges from about 0.05 mg to about 500 mg/day, e.g., from about 1 mg to about 200 mg/day, or, e.g., from about 5 mg to 25 about 175 mg/day for a 70-kg subject as loading and maintenance doses, depending on the weight of the subject, the condition and the severity of the condition.

It must be kept in mind that the materials of the present invention may generally be employed in serious disease or injury states, that is, life threatening or potentially life threatening situations. In such cases, in view of the minimization of extraneous substances and 30 general lack of immunogenicity of factor VIIa and protein C in humans, it is possible and may be felt desirable by the treating physician to administer a substantial excess of these compositions.

A dose may be given continuously or intermittently (once or several times a day). A course of treatment is typically from 1 to 30 days. In making compositions of the invention the essential ingredients; TF antagonist and Protein C are co-present and may be mixed in any 35 homogeneous or non-homogeneous manner or adjacently or otherwise proximately placed together in an individual dosage unit suitable for practicing the method of the invention. The dosage unit of the TF antagonist will usually be admixed with a carrier or inert ingredients, or

diluted by a carrier, or enclosed within a carrier which may be in the form of an ampoule, capsule, time release dosing device, sachet, paper or other container. When the carrier serves as a diluent, it may be a solid, semi-solid, paste, or liquid material which acts as a vehicle, or can be in the form of tablets, pills, powders, lozenges, elixirs, suspensions, emulsions, solutions, syrups, 5 aerosols (as a solid or in a liquid medium), or ointment, containing, for example, up to 10% by weight of the active compound. The dosage unit of the Protein C polypeptide will usually be admixed with a liquid carrier and/or other inert ingredients or enclosed within a carrier which may be in the form of an ampoule, bottle, time release dosing device or other container. When the carrier serves as a diluent, it may be a liquid material which acts as a vehicle, or can be in the 10 form of solutions containing, for example, up to 10% by weight of the active compound. The Protein C ingredient should be in an injectable liquid form immediately prior to use, however, it may be made in a storable form which is not a liquid but is easily convertible to a liquid (e.g., paste, liquid adsorbed on a solid, etc.). For the pharmaceutical formulations containing both (a) TF antagonist and (b) Protein C the carrier may be an injectable liquid medium such as is well 15 known in the art. The injectable liquid must be such that permits parenteral administration, that is, introduction of substances to a mammal being treated by interavenous, intravenous, subcutaneous, intramuscular, or intramedullary injection. Intravenous injection is most preferred as a means of administration. The Active ingredient can be dissolved or suspended in a pharmaceutically acceptable carrier, such as sterile water, sterile water containing saline and/or 20 sugars and/or suspension agents or a mixture of both. For example, for intravenous injection the compounds of the invention may be dissolved in at a concentration of about 2 mg/ml in a 4% dextrose/0.5% Na citrate aqueous solution. Liquid compositions for oral administration include pharmaceutically-acceptable emulsions, solutions, suspensions, syrups, and elixirs containing inert diluents commonly used in the art such as distilled water or ethanol. Besides inert diluents 25 such compositions may also comprise adjuvants such as wetting and suspending agents, and sweetening, flavoring, perfuming and preserving agents. Other compositions for oral administration include spray compositions which may be prepared by known methods and which comprise one or more of the active compound(s). Besides inert diluents such compositions may also comprise stabilizers such as sodium bisulfite and buffer for isotonicity, for example sodium 30 chloride, sodium citrate, or citric acid. The manufacturing of spray compositions for inhalation therapy is described in detail, for example, in the specifications of U.S. Pat. No. 2,868,691 and U.S. Pat. No. 3,095,355. Preparations for injection according to the present invention for parenteral administration include sterile aqueous or non-aqueous solutions, suspensions, or emulsions. Examples of aqueous solvents or suspending media include distilled water for 35 injection and physiological salt solution. Examples of non-aqueous solvents or suspending media are propylene glycol, polyethylene glycol, vegetable oils such as olive oil, alcohols such as ethanol, Polysorbate 80 (registered Trade Mark). These compositions may also include adjuvants

such as preserving, wetting, emulsifying and dispersing agents stabilizing agents (e.g. lactose) and solubilizers (e.g. glutamic acid and asparagine acid). They may be sterilized, for example, by filtration through a bacteria-retaining filter, by incorporation of sterilizing agents in the compositions or by irradiation. They may also be manufactured in the form of sterile solid compositions which can be dissolved in sterile water or some other sterile injectable medium immediately before use. The TF antagonist (when separate from the Protein C polypeptide) may be in the form of powder, tablet, or capsule. A solid carrier can be one or more substances which may also act as flavoring agents, lubricants, solubilizers, suspending agents, binders, tablet disintegrating agents, and encapsulating material. Suitable solid carriers are magnesium carbonate, magnesium stearate, talc, sugar lactose, pectin, dextrin, starch, gelatin, tragacanth, methyl cellulose, sodium carboxymethyl cellulose, low melting waxes, and cocoa butter. Preferably, the pharmaceutical compositions are administered parenterally, i.e., intravenously, subcutaneously, or intramuscularly; intravenously being most preferred. They may also be administered by continuous or pulsatile infusion. Local delivery of the preparations of the present invention, such as, for example, topical application, may be carried out, e.g., by means of a spray, perfusion, double balloon catheters, stent, incorporated into vascular grafts or stents, hydrogels used to coat balloon catheters, or other well established methods. One skilled in this art may formulate the compositions of the invention an appropriate manner, and in accordance with accepted practices, such as those disclosed in Remington's Pharmaceutical Sciences, 20 Gennaro, ed., Mack Publishing Co., Easton, PA, 1990.

This invention is a method of treating or preventing thrombotic and coagulopathic diseases, Inflammatory Disease or Respiratory Disease by administering to a mammal in need thereof a therapeutically effective amount of (a) a TF antagonist and b) a Protein C polypeptide; wherein (a) and (b) are both administered within a therapeutically effective interval. The 25 administration of (a) or (b) to, e.g., a septic patient may be either continuous or intermittent.

The Protein C polypeptide and a TF antagonist can be delivered simultaneously. One convenient method of simultaneous delivery is to use the compositions of the invention, wherein the Active Ingredient has the essential ingredients co-present in a unit dosage form. Solutions or suspensions of mixed essential ingredients may, if desired, be delivered from the same liquid 30 holding bag. Another method of simultaneous delivery of the Protein C polypeptide and a TF antagonist is to deliver them to the patient separately but simultaneously. Thus, for example, some TF antagonists may be given as an oral formulation at the same time as the Protein C polypeptide is given parenterally. Dosage of a TF antagonist can begin simultaneously with the Protein C administration. The length of the TF antagonist administration can extend past the 35 Protein C administration, or vice versa.

Each of the essential ingredients, viz., a therapeutically effective amount of (a) a TF antagonist in and (b) Protein C polypeptide have a therapeutically effective interval, namely the

interval of time in which each agent provides benefit for the patient being treated with Thrombotic or Coagulopathic related disease, Inflammatory Disease or Respiratory Disease. The method of the invention may be practiced by separately dosing the patient in any order with a therapeutically effective amount of (a) a TF antagonist and (b) Protein C polypeptide provided that each agent is given within the period of time that that the other agent is therapeutically effective against Thrombotic or Coagulopathic related disease, Inflammatory Disease or Respiratory Disease, or organ failure resulting from these pathologic processes.

5 The Protein C polypeptide and TF antagonist are preferably administered parenterally to a patient to insure their delivery into the bloodstream in an effective form as fast as possible.

10 The amount and relative ratio of protein C polypeptide and TF antagonist to be used in the practice of the method of invention is set out in the previous section. It may be appreciated that it may be necessary to make routine variations to the dosage of either agent depending on the age and condition of the patient. The decision to begin the therapy will be based upon the appearance of the clinical manifestations of Thrombotic or Coagulopathic related disease,

15 Inflammatory Disease or Respiratory Disease. Typical clinical manifestations are coughing, restricted breathing, obstructed airway, pain, fever, chills, tachycardia, tachypnea, altered mental state, hypothermia, hyperthermia, accelerated or repressed breathing or heart rates, increased or decreased white blood cell count, and hypotension. For Respiratory Disease diagnostic tests such as roetgenographic examination, bronchoscopy, lung biopsy, spirography (lung capacity, residual 20 volume, flow rates, etc.) are used. These and other symptoms and diagnostic techniques are well known in the art as set out in standard references such as, Harrison's Principles of Internal medicine (ISBN 0-07-032370-4) 1994.

The decision to determine the length of therapy may be supported by standard clinical laboratory results from commercially available assays or instrumentation supporting the 25 eradication of the symptoms defining Thrombotic or Coagulopathic related disease, Inflammatory or Respiratory Diseases. The method of the invention may be practiced by continuously or intermittently administering a therapeutically effective dose of the essential Protein C and TF antagonist ingredients for as long as deemed efficacious for the treatment of the episode. The administration can be conducted for up to a total of about 60 days with a 30 preferred course of therapy lasting for up to 14 days. The therapy may be restarted upon the return of the Thrombotic or Coagulopathic related disease, Inflammatory or Respiratory disease.

While the present invention has been illustrated above by certain specific embodiments, it is not intended that these specific examples should limit the scope of the invention as described in the appended claims.

**Assays:****Factor VII biological activity**

The biological activity of factor VIIa in blood clotting derives from its ability to (i) bind to tissue factor (TF) and (ii) catalyze the proteolytic cleavage of factor IX or factor X to produce activated factor IX or X (factor IXa or Xa, respectively).

5 Biological activity of factor VII polypeptides ("factor VII biological activity") may be quantified by measuring the ability of a preparation to promote blood clotting using factor VII-deficient plasma and thromboplastin, as described, e.g., in U.S. Patent No. 5,997,864. In this assay, biological activity is expressed as the reduction in clotting time relative to a control sample and is converted to "factor VII units" by comparison with a pooled human serum standard containing 1 unit/ml factor VII activity. Alternatively, factor VIIa biological activity may be  
10 quantified by

- (i) Measuring the ability of factor VIIa or a factor VIIa -related polypeptide to produce activated factor X (factor Xa) in a system comprising TF embedded in a lipid membrane and factor X. (Persson et al., J. Biol. Chem. 272:19919-19924, 1997);
- (ii) Measuring factor X hydrolysis in an aqueous system ("In Vitro Proteolysis Assay", see  
15 below);
- (iii) Measuring the physical binding of factor VIIa or a factor VIIa -related polypeptide to TF using an instrument based on surface plasmon resonance (Persson, FEBS Letts. 413:359-363, 1997); and
- (iv) Measuring hydrolysis of a synthetic substrate by factor VIIa and/or a factor VIIa -related  
20 polypeptide ("In Vitro Hydrolysis Assay", see below); and
- (v) Measuring generation of thrombin in a TF-independent in vitro system.

In Vitro Hydrolysis Assay

Native (wild-type) factor VIIa and factor VIIa variant (both hereafter referred to as  
25 "factor VIIa") may be assayed for specific activities. They may also be assayed in parallel to directly compare their specific activities. The assay is carried out in a microtiter plate (MaxiSorp, Nunc, Denmark). The chromogenic substrate D-Ile-Pro-Arg-p-nitroanilide (S-2288, Chromogenix, Sweden), final concentration 1 mM, is added to factor VIIa (final concentration 100 nM) in 50 mM Hepes, pH 7.4, containing 0.1 M NaCl, 5 mM CaCl<sub>2</sub> and 1 mg/ml bovine serum albumin. The  
30 absorbance at 405 nm is measured continuously in a SpectraMax™ 340 plate reader (Molecular Devices, USA). The absorbance developed during a 20-minute incubation, after subtraction of the absorbance in a blank well containing no enzyme, is used to calculate the ratio between the activities of variant and wild-type factor VIIa:

$$\text{Ratio} = (A_{405 \text{ nm}} \text{ factor VIIa variant}) / (A_{405 \text{ nm}} \text{ factor VIIa wild-type}).$$

35 Based thereon, factor VIIa variants with an activity comparable to or higher than native factor VIIa may be identified, such as, for example, variants where the ratio between the activity of the variant and the activity of native factor VII (wild-type FVII) is around, versus above 1.0.

The activity of factor VIIa or factor VIIa variants may also be measured using a physiological substrate such as factor X, suitably at a concentration of 100-1000 nM, where the factor Xa generated is measured after the addition of a suitable chromogenic substrate (e.g. S-2765). In addition, the activity assay may be run at physiological temperature.

5

In Vitro Proteolysis Assay

Native (wild-type) factor VIIa and factor VIIa variant (both hereafter referred to as "factor VIIa") are assayed in parallel to directly compare their specific activities. The assay is carried out in a microtiter plate (MaxiSorp, Nunc, Denmark). factor VIIa (10 nM) and factor X 10 (0.8 microM) in 100 microL 50 mM Hepes, pH 7.4, containing 0.1 M NaCl, 5 mM CaCl2 and 1 mg/ml bovine serum albumin, are incubated for 15 min. factor X cleavage is then stopped by the addition of 50 microL 50 mM Hepes, pH 7.4, containing 0.1 M NaCl, 20 mM EDTA and 1 mg/ml bovine serum albumin. The amount of factor Xa generated is measured by addition of the 15 chromogenic substrate Z-D-Arg-Gly-Arg-p-nitroanilide (S-2765, Chromogenix, Sweden), final concentration 0.5 mM. The absorbance at 405 nm is measured continuously in a SpectraMax™ 340 plate reader (Molecular Devices, USA). The absorbance developed during 10 minutes, after subtraction of the absorbance in a blank well containing no FVIIa, is used to calculate the ratio between the proteolytic activities of variant and wild-type factor VIIa:

$$\text{Ratio} = (\text{A405 nm factor VIIa variant})/(\text{A405 nm factor VIIa wild-type}).$$

20 Based thereon, factor VIIa variants with an activity comparable to or higher than native factor VIIa may be identified, such as, for example, variants where the ratio between the activity of the variant and the activity of native factor VII (wild-type FVII) is around, versus above 1.0.

Inhibition of FVIIa/phospholipids-embedded TF-catalyzed activation of FX by TF antagonists –

25 FXa generation assay (assay 1):

In the following example all concentrations are final. Lipidated TF (10 pM), FVIIa (100 pM) and TF antagonist or FFR-rFVIIa (0 – 50 nM) in HBS/BSA (50 mM hepes, pH 7.4, 150 mM NaCl, 5 mM CaCl2, 1 mg/ml BSA) are incubated 60 min at room temperature before FX (50 nM) is added. The reaction is stopped after another 10 min by addition of ½ volume stopping buffer 30 (50 mM Hepes, pH 7.4, 100 mM NaCl, 20 mM EDTA). The amount of FXa generated is determined by adding substrate S2765 (0.6 mM, Chromogenix, and measuring absorbance at 405 nm continuously for 10 min. IC50 values for TF antagonist inhibition of FVIIa/lipidated TF-mediated activation of FX may be calculated. The IC50 value for FFR-rFVIIa is 51 +/- 26 pM in this assay.

35

Inhibition of FVIIa/cell surface TF-catalyzed activation of FX by TF antagonists –

FXa generation assay (Assay 2):

In the following example all concentrations are final. Monolayers of human lung fibroblasts WI-38 (ATTC No. CCL-75) or human bladder carcinoma cell line J82 (ATTC No. HTB-1) or human keratinocyte cell line CCD 1102KerTr (ATCC no. CRL-2310) constitutively expressing TF are employed as TF source in FVIIa/TF catalyzed activation of FX. Confluent cell monolayers in a 5 96-well plate are washed one time in buffer A (10 mM Hepes, pH 7.45, 150 mM NaCl, 4 mM KCl, and 11 mM glucose) and one time in buffer B (buffer A supplemented with 1 mg/ml BSA and 5 mM Ca<sup>2+</sup>), FVIIa (1 nM), FX (135 nM) and varying concentrations of TF antagonist or FFR-rFVIIa in buffer B are simultaneously added to the cells. FXa formation is allowed for 15 min at 37°C. 50-  
10 µl aliquots are removed from each well and added to 50 µl stopping buffer (Buffer A supplemented with 10 mM EDTA and 1 mg/ml BSA). The amount of FXa generated is determined by transferring 50 µl of the above mixture to a microtiter plate well and adding 25 µl Chromozym X (final concentration 0.6 mM) to the wells. The absorbance at 405 nm is measured continuously and the initial rates of color development are converted to FXa concentrations using an FXa standard curve. The IC<sub>50</sub> value for FFR-rFVIIa is 1.5 nM in this assay.

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Inhibition of 125I-FVIIa binding to cell surface TF by TF antagonists –

TF binding assay (Assay 3):

In the following example all concentrations are final. Binding studies are employed using the human bladder carcinoma cell line J82 (ATTC No. HTB-1) or the human keratinocyte 20 cell line (CCD1102KerTr ATCC No CRL-2310) or NHEK P166 (Clonetics No. CC-2507) all constitutively expressing TF. Confluent monolayers in 24-well tissue culture plates are washed once with buffer A (10 mM Hepes, pH 7.45, 150 mM NaCl, 4 mM KCl, and 11 mM glucose) supplemented with 5 mM EDTA and then once with buffer A and once with buffer B (buffer A supplemented with 1 mg/ml BSA and 5 mM Ca<sup>2+</sup>). The monolayers are preincubated 2 min with 25 100 µl cold buffer B. Varying concentrations of Mabs (or FFR-FVIIa) and radiolabelled FVIIa (0.5 nM 125I-FVIIa) are simultaneously added to the cells (final volume 200 µl). The plates are incubated for 2 hours at 4 °C. At the end of the incubation, the unbound material is removed, and the cells are washed 4 times with ice-cold buffer B and lysed with 300 µl lysis buffer (200 mM NaOH, 1 % SDS and 10 mM EDTA). Radioactivity is measured in a gamma counter (Cobra, 30 Packard Instruments). The binding data are analyzed and curve fitted using GraFit4 (Eritacus Software, Ltd., (U.K.). The IC<sub>50</sub> value for FFR-rFVIIa is 4 nM in this assay.

Biosensor assay (Assay 4):

TF antagonists are tested on the Biacore instrument by passing a standard solution of 35 the TF antagonist over a chip with immobilized TF. This is followed by different concentrations of sTF in 10 mM hepes pH 7.4 containing 150 mM NaCl, 10 mM CaCl<sub>2</sub> and 0.0003 % polysorbate 20. Kds are calculated from the sensorgrams using the integrated Biacore evaluation software.

Test for protein C activity:

A suitable assay for testing for Protein C anticoagulant and amidolytic activity and thereby selecting suitable protein C variants can be performed as described, for example, in 5 Grinnell et al., 1987, Bio/Technology 5:1189-1192 ("the Protein C Assay"), which is hereby incorporated by reference in its entirety.

The present invention is further illustrated by the following examples, which, however, are not to be construed as limiting the scope of protection. The features disclosed in the foregoing description and in the following examples may, both separately and in any combination 10 thereof, be material for realizing the invention in diverse forms thereof.

#### EXAMPLES

a. Preparation of Human Protein C

15 Recombinant human Protein C (r-hPC) was produced in Human Kidney 293 cells by techniques well known to the skilled artisan such as those set forth in Yan, U.S. Patent No. 4,981,952, the entire disclosure of which is herein incorporated by reference. The gene encoding human Protein C is disclosed and claimed in Bang et al., U.S. Patent No. 4,775,624, the entire disclosure of which is incorporated herein by reference. The plasmid used to express human 20 Protein C in 293 cells was plasmid pLPC which is disclosed in Bang et al., U.S. Patent No. 4,992,373, the entire disclosure of which is incorporated herein by reference. The construction of plasmid pLPC is also described in European Patent Publication No. 0 445 939, the teachings of which are also incorporated herein by reference and in Grinnell et al., 1987, Bio/Technology 5:1189-1192. Briefly, the plasmid was transfected into 293 cells, and thereafter stable 25 transforms were identified, subcultured, and grown in serum-free media. After fermentation, cell-free medium was obtained by microfiltration.

The human Protein C was separated from the culture fluid by an adaptation of the techniques of Yan, U.S. Patent No. 4,981,952, the entire disclosure of which is herein incorporated by reference. The clarified medium was made 4 mM in EDTA before it was 30 absorbed to an anion exchange resin (Fast-Flow Q, Pharmacia). After washing with 4 column volumes of 20 mM Tris, 200 mM NaCl, pH 7.4 and 2 column volumes of 20 mM Tris, 150 mM NaCl, pH 7.4, the bound recombinant human Protein C zymogen was eluted with 20 mM Tris, 150 mM NaCl, 10 mM CaCl<sub>2</sub>, pH 7.4. The eluted protein was greater than 95% pure after elution as judged by SDS-polyacrylamide gel electrophoresis. Further purification of the protein was 35 accomplished by making the protein 3 M in NaCl followed by adsorption to a hydrophobic interaction resin (Toyopearl Phenyl 650M, TosoHaas) equilibrated in 20 mM Tris, 3 M NaCl, 10 mM CaCl<sub>2</sub>, pH 7.4. After washing with 2 column volumes of equilibration buffer without CaCl<sub>2</sub>,

the recombinant human Protein C was eluted with 20 mM Tris, pH 7.4. The eluted protein was prepared for activation by removal of residual calcium. The recombinant human Protein C was passed over a metal affinity column (Chelex-100, Bio-Rad) to remove calcium and again bound to an anion exchanger (Fast Flow Q, Pharmacia). Both of these columns were arranged in series  
5 and equilibrated in 20 mM Tris, 1'50 mM NaCl, 5 mM EDTA, pH 7.4. Following loading of the protein, the Chelex-100 column was washed with one column volume of the same buffer before disconnecting it from the series. The anion exchange column was washed with 3 column volumes of equilibration buffer before eluting the protein with 0.4 M NaCl, 20 mM Tris-acetate, pH 6.5. Protein concentrations of recombinant human protein C and recombinant activated  
10 Protein C solutions were measured by UV 280 nm extinction  $E_{0.1\%}=1.85$  or 1.95, respectively.

b. Activation of recombinant human Protein C

Bovine thrombin was coupled to Activated CH-Sepharose 4B (Pharmacia) in the presence of 50 mM HEPES, pH 7.5 at 4 OC. The coupling reaction was done on resin already packed into a column using approximately 5000 units thrombin/ml resin. The thrombin solution 5 was circulated through the column for approximately 3 hours before adding MEA to a concentration of 0.6 ml/l of circulating solution. The MEA-containing solution was circulated for an additional 10-12 hours to assure complete blockage of the unreacted amines on the resin. Following blocking, the thrombin-coupled resin was washed with 10 column volumes of 1 M NaCl, 20 mM Tris, pH 6.5 to remove all non-specifically bound protein, and was used in activation 10 reactions after equilibrating in activation buffer. Purified rHPC was made 5 mM in EDTA (to chelate any residual calcium) and diluted to a concentration of 2 mg/ml with 20 mM Tris, pH 7.4 or 20 mM Tris-acetate, pH 6.5. This material was passed through a thrombin column equilibrated at 37(C with 50 mM NaCl and either 20 mM Tris pH 7.4 or 20 mM Tris-acetate pH 6.5. The flow rate was adjusted to allow for approximately 20 min. of contact time between the rHPC and 15 thrombin resin. The effluent was collected and immediately assayed for amidolytic activity. If the material did not have a specific activity (amidolytic) comparable to an established standard of aPC, it was recycled over the thrombin column to activate the rHPC to completion. This was followed by 1:1 dilution of the material with 20 mM buffer as above, with a pH of either 7.4 or 6.5 to keep the aPC at lower concentrations while it awaited the next processing step. Removal 20 of leached thrombin from the aPC material was accomplished by binding the aPC to an anion exchange resin (Fast Flow Q, Pharmacia) equilibrated in activation buffer (either 20 mM Tris, pH 7.4 or 20 mM Tris-acetate, pH 6.5) with 150 mM NaCl. Thrombin does not interact with the anion exchange resin under these conditions, but passes through the column into the sample application effluent. Once the aPC is loaded onto the column, a 2-6 column volume wash with 25 20 mM equilibration buffer is done before eluting the bound aPC with a step elution using 0.4 M NaCl in either 5 MM Tris-acetate, pH 6.5 or mM Tris, pH 7.4. Higher volume washes of the column facilitated more complete removal of the dodecapeptide. The material eluted from this column was stored either in a frozen solution (-20 OC) or as a lyophilized powder. The anticoagulant activity of activated Protein C was determined by measuring the prolongation 30 of the clotting time in the activated partial thromboplastin time (APTT) clotting assay. A standard curve was prepared in dilution buffer (1 mg/ml radioimmunoassay grade BSA, 20 mM Tris, pH 7.4, 150 mM NaCl, 0.02% NaN3) ranging in Protein C concentration from 125 -1000 ng/ml. Samples were prepared at several dilutions in this concentration range. To each sample cuvette, 50 gl of cold horse plasma and 50 gl of reconstituted activated partial thromboplastin 35 time reagent (APTT Reagent, Sigma) were added and incubated at 37 OC for 5 min. After incubation, 50 gl of the appropriate samples or standards were added to each cuvette. Dilution buffer was used in place of sample or standard to determine basal clotting time. The timer of

the fibrometer (CoA Screener Hemostasis Analyzer, American Laboratory) was started immediately after the addition of 50 gl 37 (C 30 mM CaC12 to each sample or standard. Activated Protein C concentration in samples are calculated from the linear regression equation of the standard curve. Clotting times reported here are the average of a minimum of three replicates, including standard curve samples.